



# **STIC Search Report**

## **EIC 1700**

**STIC Database Tracking Number: 174287**

**TO: Gregg Cantelmo**  
**Location: REM 6B71**  
**Art Unit : 1745**  
**December 21, 2005**

**Case Serial Number: 10/620687**

**From: Ross Shipe**  
**Location: EIC 1700**  
**REMSSEN 4B31**  
**Phone: 571/272-6018**  
**Ross.Shipe@uspto.gov**

### **Search Notes**

Examiner Cantelmo:

Please review the attached search results.

If you have any questions or if you would like to refine the search query, please feel free to contact me at any time.

Thanks you for using EIC 1700 search services!

Ross Shipe (ASRC)  
Technical Information Specialist

**SEARCH REQUEST FORM****Scientific and Technical Information Center**

Requester's Full Name: GREG CASTELMO Examiner #: \_\_\_\_\_ Date: \_\_\_\_\_  
 Art Unit: \_\_\_\_\_ Phone Number 30 \_\_\_\_\_ Serial Number: 10/620652  
 Mail Box and Bldg/Room Location: \_\_\_\_\_ Results Format Preferred (circle): PAPER DISK E-MAIL

**If more than one search is submitted, please prioritize searches in order of need.**

\*\*\*\*\*

Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc, if known. Please attach a copy of the cover sheet, pertinent claims, and abstract.

Title of Invention: Lithium Rechargeable Battery  
 Inventors (please provide full names): \_\_\_\_\_

Earliest Priority Filing Date: \_\_\_\_\_

*\*For Sequence Searches Only\* Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.*

\*\*\*\*\*  
**STAFF USE ONLY**

	Type of Search	Vendors and cost where applicable
Searcher: <u>ROS</u>	NA Sequence (#) _____	STN <u>✓</u>
Searcher Phone #: _____	AA Sequence (#) _____	Dialog _____
Searcher Location: _____	Structure (#) _____	Questel/Orbit _____
Date Searcher Picked Up: _____	Bibliographic <u>✓</u>	Dr. Link _____
Date Completed: <u>12/21/05</u>	Litigation _____	Lexis/Nexis _____
Searcher Prep & Review Time: <u>30</u>	Fulltext _____	Sequence Systems _____
Clerical Prep Time: _____	Patent Family _____	WWW/Internet _____
Online Time: <u>30 108</u>	Other _____	Other (specify) _____

**Mellerson, Kendra**

174287

**From:** "Gregg Cantelmo" [gregg.cantelmo]  
**Sent:** Wednesday, December 14, 2005 4:05 PM  
**To:** STIC-EIC1700  
**Subject:** Database Search Request, Serial Number: 10/620687

**Requester:**  
Gregg Cantelmo (TC1700)  
**Art Unit:**  
1745  
**Employee Number:**  
75777  
**Office Location:**  
REM 6B71  
**Phone Number:**  
571-272-1283  
**Mailbox Number:**

**Case serial number:**  
10/620687  
**Class / Subclass(es):**

**Earliest Priority Filing Date:**

**Format preferred for results:**  
Paper

**Search Topic Information:**  
Alleged novelty lies in the mixed graphite material of claim 1/  
**Special Instructions and Other Comments:**

SCIENTIFIC REFERENCE BR  
Sci & Tech Inf. Cntr

DEC 15 RECD

Pat. & T.M. Office



# STIC Search Results Feedback Form

**EIC17000**

Questions about the scope or the results of the search? Contact *the EIC searcher* or contact:

Kathleen Fuller, EIC 1700 Team Leader  
571/272-2505 REMSEN 4B28

## Voluntary Results Feedback Form

- I am an examiner in Workgroup:  Example: 1713
- Relevant prior art **found**, search results used as follows:

- ☐ 102 rejection
- ☐ 103 rejection
- ☐ Cited as being of interest.
- ☐ Helped examiner better understand the invention.
- ☐ Helped examiner better understand the state of the art in their technology.

Types of relevant prior art found:

- ☐ Foreign Patent(s)
- ☐ Non-Patent Literature  
(journal articles, conference proceedings, new product announcements etc.)

- Relevant prior art **not found**:

- ☐ Results verified the lack of relevant prior art (helped determine patentability).
- ☐ Results were not useful in determining patentability or understanding the invention.

Comments:

=> d his full

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FILE 'REGISTRY' ENTERED AT 09:48:46 ON 21 DEC 2005
L1      1 SEA ABB=ON  PLU=ON  COPPER/CN
L2      1 SEA ABB=ON  PLU=ON  GRAPHITE/CN

FILE 'HCAPLUS' ENTERED AT 09:49:13 ON 21 DEC 2005
L3      1169095 SEA ABB=ON  PLU=ON  COPPER OR CU OR L1
L4      715431 SEA ABB=ON  PLU=ON  ELECTRODE# OR CATHODE#
L5      191171 SEA ABB=ON  PLU=ON  GRAPHITE# OR BLACK (2A) LEAD OR
      PLUMBAGO## OR L2
L6      2533705 SEA ABB=ON  PLU=ON  ARTIFICIAL? OR SYNTHETIC? OR
      SIMULAT? OR FABRICAT? OR MANMADE OR MAN (A) MADE OR
      ERSATZ OR SUBSTITUT?
L7      827599 SEA ABB=ON  PLU=ON  SPHERICAL? OR ROUND? OR CIRCULAR? OR
      CURV? OR SPHERIOD##
L9      6 SEA ABB=ON  PLU=ON  L7 (2A) L5 AND L6 (2A) L5 AND L4
L12     133 SEA ABB=ON  PLU=ON  L7 (L) L5 AND L6 (L) L5 (L) L4
L13     18855 SEA ABB=ON  PLU=ON  MESOPHASE OR MESOPHASIC
L14     48953 SEA ABB=ON  PLU=ON  PITCH##
L16     2 SEA ABB=ON  PLU=ON  L13 AND L12
L19     6 SEA ABB=ON  PLU=ON  L13 AND L14 AND L4 AND L5 AND L3
L20     51 SEA ABB=ON  PLU=ON  L13 (L) L14 (L) L4 (L) L5
L21     24 SEA ABB=ON  PLU=ON  L13 (L) L14 (L) L4 (L) L5 AND
      ELECTRO?/SC,SC
L22     27 SEA ABB=ON  PLU=ON  L20 NOT L21
L23     25 SEA ABB=ON  PLU=ON  L22 AND (57? OR 51?)/SC
L26     25 SEA ABB=ON  PLU=ON  L23 NOT (L9 OR L16 OR L19 OR L21)
L28     10 SEA ABB=ON  PLU=ON  L26 AND (ELECTRODE# OR CATHODE#)/TI
L29     43 SEA ABB=ON  PLU=ON  L9 OR L16 OR L19 OR L21 OR L28

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FILE 'WPIX' ENTERED AT 11:08:57 ON 21 DEC 2005
L30     9 SEA ABB=ON  PLU=ON  L9 OR L16 OR L19 OR L21 OR L28

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FILE 'INSPEC' ENTERED AT 11:11:39 ON 21 DEC 2005
L31     4 SEA ABB=ON  PLU=ON  L9 OR L16 OR L19 OR L21 OR L28

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FILE 'JICST-EPLUS' ENTERED AT 11:16:02 ON 21 DEC 2005
L32     0 SEA ABB=ON  PLU=ON  L9 OR L16 OR L19 OR L21 OR L28

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FILE 'JAPIO' ENTERED AT 11:16:31 ON 21 DEC 2005
L33     6 SEA ABB=ON  PLU=ON  L9 OR L16 OR L19 OR L21 OR L28

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FILE 'COMPENDEX' ENTERED AT 11:17:21 ON 21 DEC 2005
L34     3 SEA ABB=ON  PLU=ON  L9 OR L16 OR L19 OR L21 OR L28

```

=> file wpix

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FILE 'WPIX' ENTERED AT 11:21:03 ON 21 DEC 2005
COPYRIGHT (C) 2005 THE THOMSON CORPORATION

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=> d que l30

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L1      1 SEA FILE=REGISTRY ABB=ON  PLU=ON  COPPER/CN
L2      1 SEA FILE=REGISTRY ABB=ON  PLU=ON  GRAPHITE/CN
L3      1169095 SEA FILE=HCAPLUS ABB=ON  PLU=ON  COPPER OR CU OR L1
L4      715431 SEA FILE=HCAPLUS ABB=ON  PLU=ON  ELECTRODE# OR CATHODE#
L5      191171 SEA FILE=HCAPLUS ABB=ON  PLU=ON  GRAPHITE# OR BLACK (2A)
      LEAD OR PLUMBAGO## OR L2
L6      2533705 SEA FILE=HCAPLUS ABB=ON  PLU=ON  ARTIFICIAL? OR SYNTHETIC
      ? OR SIMULAT? OR FABRICAT? OR MANMADE OR MAN (A) MADE OR

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L7 827599 ERSATZ OR SUBSTITUT?  
 SEA FILE=HCAPLUS ABB=ON PLU=ON SPHERICAL? OR ROUND? OR  
 CIRCULAR? OR CURV? OR SPHERIOD##  
 L9 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L7 (2A) L5 AND L6 (2A)  
 L5 AND L4  
 L12 133 SEA FILE=HCAPLUS ABB=ON PLU=ON L7 (L) L5 AND L6 (L) L5  
 (L) L4  
 L13 18855 SEA FILE=HCAPLUS ABB=ON PLU=ON MESOPHASE OR MESOPHASIC  
 L14 48953 SEA FILE=HCAPLUS ABB=ON PLU=ON PITCH##  
 L16 2 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 AND L12  
 L19 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 AND L14 AND L4 AND  
 L5 AND L3  
 L20 51 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 (L) L14 (L) L4 (L)  
 L5  
 L21 24 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 (L) L14 (L) L4 (L)  
 L5 AND ELECTRO?/SC,SC  
 L22 27 SEA FILE=HCAPLUS ABB=ON PLU=ON L20 NOT L21  
 L23 25 SEA FILE=HCAPLUS ABB=ON PLU=ON L22 AND (57? OR 51?)/SC  
 L26 25 SEA FILE=HCAPLUS ABB=ON PLU=ON L23 NOT (L9 OR L16 OR  
 L19 OR L21)  
 L28 10 SEA FILE=HCAPLUS ABB=ON PLU=ON L26 AND (ELECTRODE# OR  
 CATHODE#)/TI  
 L30 9 SEA FILE=WPIX ABB=ON PLU=ON L9 OR L16 OR L19 OR L21 OR  
 L28

=> d 130 full 1-9

L30 ANSWER 1 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN  
 AN 2005-435450 [44] WPIX  
 DNN N2005-353359 DNC C2005-133644  
 TI Production of carbon negative electrode materials for  
 lithium-ion batteries, for applications in electrical and electronic  
 industries particularly for e.g. mobile phones, portable computers  
 and digital cameras.  
 DC E36 L03 X16  
 IN HE, Z; MA, J; WANG, J; DING, X; FENG, S; FU, Z; LI, S; ZHANG, D  
 PA (SHAN-N) SHANGHAI SHANSHAN SCI & TECHNOLOGY CO LTD; (SHAN-N)  
 SHANGHAI SHANSHAN TECH CO LTD  
 CYC 108  
 PI WO 2005055346 A1 20050616 (200544)\* ZH 13 H01M004-58  
 RW: AT BE BG BW CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE  
 IS IT KE LS LU MC MW MZ NA NL OA PL PT RO SD SE SI SK SL SZ  
 TR TZ UG ZM ZW  
 W: AE AG AL AM AT AU AZ BA BB BG BR BW BY BZ CA CH CN CO CR CU  
 CZ DE DK DM DZ EC EE EG ES FI GB GD GE GH GM HR HU ID IL IN  
 IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW  
 MX MZ NA NI NO NZ OM PG PH PL PT RO RU SC SD SE SG SK SL SY  
 TJ TM TN TR TT TZ UA UG US UZ VC VN YU ZA ZM ZW  
 CN 1624955 A 20050608 (200566) H01M004-04  
 ADT WO 2005055346 A1 WO 2004-CN1297 20041115; CN 1624955 A CN  
 2003-1108982 20031201  
 PRAI CN 2003-1108982 20031201  
 IC ICM H01M004-04; H01M004-58  
 ICS C01B031-00  
 AB WO2005055346 A UPAB: 20050712  
 NOVELTY - A process for producing carbon negative electrode  
 materials for lithium-ion batteries comprises supplying a coating  
 material of natural graphite and a coating material of heavy  
 aromatic hydrocarbon or its mixture, polymerization to give a

coating layer with microcapsulation on surface of the natural graphite, extraction and separation then drying to a dried product for carbonizing or graphitizing.

DETAILED DESCRIPTION - A process for producing carbon negative electrode materials for lithium-ion batteries comprises:

(1) supplying a coating material (A) of natural graphite and a coating material (B) of heavy aromatic hydrocarbon or its mixture, in a weight ratio of (A):(B) = 1:0.5-10;

(2) polymerization of the resultant solid-liquid mixture at 350-500 deg. C, at a pressure of 0.01-10 MPa for 5-450 minutes to give a coating layer with microcapsulation on surface of the natural graphite;

(3) extraction and separation of solid-phase granules in the reaction system;

(4) drying to remove the light component in such solid-phase granules to a dried product that has a weight gain of 2-100%, with respect to graphite raw material; and

(5) carbonizing or graphitizing the product to obtain a carbon negative electrode material that has a surface-coated layer of artificial carbon or artificial graphite at 80-220 deg. C or 2400-3000 deg. C, respectively.

USE - The thus produced batteries are applications in electrical and electronic industries particularly for e.g. mobile phones, portable computers and digital cameras.

ADVANTAGE - Such lithium-ion batteries thus made have high specific capacity and long cycle lifetime.

Dwg.0/0

TECH WO 2005055346 A1UPTX: 20050712

TECHNOLOGY FOCUS - INORGANIC CHEMISTRY - Preferred Materials: The natural graphite is particularly spherical or plate-like natural graphite.

TECHNOLOGY FOCUS - ORGANIC CHEMISTRY - Preferred Materials: The heavy aromatic hydrocarbon or its mixture is selected from tar coal, coal pitch, petroleum pitch and their mixture. The method for separation is by applying a solvent, centrifugation or precipitation. Such method with solvent for separation can employ a solvent chosen from anthracene oil, washing oil, diesel oil, pyridine, quinoline, xylene and toluene, in a weight ratio of such solvent to the reaction product of 0.5-5:1. During drying, the temperature is set at 200-500degreesC. When graphitizing, the graphitization temperature is particularly at 2800- 300degreesC.

ABEX WO 2005055346 A1UPTX: 20050712

EXAMPLE - Spherical natural graphite and tar coal (1:1 in ratio) were reacted at 400degreesC and 0.1 MPa for 60 minutes. After work-up, the resulting product was carbonized at 1000degreesC. Properties of the thus obtained electrode material: D50 = 20.1 microns; weight gain = 6.2%; true density = 2.18 g/cm3; vibration-packed density = 1.25 g/cm3; specific area = 2.6 m2/g; initial discharge capacity = 335 mAh/g; and initial discharge efficiency = 94.3%.

FS CPI EPI

FA AB; DCN

MC CPI: E31-N04B; L03-E01B5

EPI: X16-E01C; X16-E03A; X16-E08

L30 ANSWER 2 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN

AN 2004-273975 [26] WPIX

DNN N2004-216708 DNC C2004-107064

TI Activated carbon, useful for electrical double layer capacitor, has preset specific surface area and pore volume.

DC E36 L03 V01 X16 X21

PA (HOND) HONDA MOTOR CO LTD; (KASH-N) KASHIMA SEKIYU KK; (KURS)  
KURARAY CHEM CO LTD

CYC 1  
PI JP 2004067498 A 20040304 (200426)\* 18 C01B031-12

ADT JP 2004067498 A JP 2003-150869 20030528

PRAI JP 2002-173146 20020613

IC ICM C01B031-12

ICS H01G009-058

AB JP2004067498 A UPAB: 20040421

NOVELTY - Activated carbon has Brunauer Emmett Teller (BET) specific surface area of less than 1200 m<sup>2</sup>/g by nitrogen absorption process, and pore volume of 0.3 ml/g or more, measured with t-plotting method by nitrogen adsorption.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

(1) manufacture of activated carbon; and

(2) electrical double layer capacitor (1).

USE - For electrical double layer capacitor (claimed).

ADVANTAGE - The activated carbon is recyclable for a long period of time, and provides an electrical double layer capacitor having high volume capacitance and durability.

DESCRIPTION OF DRAWING(S) - The figure shows a typical perspective view of the electrical double layer capacitor.

electrical double layer capacitor 1

pressure plates 3, 4

electrodes 5, 6

aluminum meshes 7, 8

aluminum wires 9, 10

Dwg.1/4

TECH JP 2004067498 AUPTX: 20040421

TECHNOLOGY FOCUS - INORGANIC CHEMISTRY - Preferred Properties: The BET specific surface area is 700-900 m<sup>2</sup>/g and the pore volume is 0.3-0.4 ml/g. The molding density of activated carbon by electrolyzing process is 0.80-0.95 g/cm<sup>3</sup>, and the electrical double layer capacitor has capacitance of more than 30 F/cm<sup>3</sup>. Preferred Composition: The activated carbon contains 4 wt.% or less of oxygen, and at least one type of functional group with respect to surface functional groups containing 0.5 meq/g or less of carboxyl, quinone, hydroxyl and lactone groups by hydrochloric acid titrimetric method. The activated carbon may further comprise 20 ppm or less of nickel, 5 ppm or less of iron, 1 ppm or less of copper and silver, and 200 ppm or less of transition metal (compound).

Preferred Carbon: The activated carbon is obtained by alkali activation processing a carbon material having graphite property.

Preferred Precursor: The carbon precursor is a synthetic mesophase pitch.

FS CPI EPI

FA AB; GI; DCN

MC CPI: E11-Q01; E11-Q02; E31-H03; E31-N03; E31-N04C; L03-B03A

EPI: V01-B01A; V01-B01D; X16-L02; X21-B04

L30 ANSWER 3 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN

AN 2004-156212 [15] WPIX

DNN N2004-125079 DNC C2004-061955

TI Lithium rechargeable battery for use as driving power source, includes negative electrode having active mixture layer of artificial and spherical graphite particles fixed on copper core material.

DC A85 L03 X16

IN FUJIWARA, S; FUKUMOTO, Y; KATO, F; OURA, T; YAMAMOTO, N

*Applicant*



PA (MATU) MATSUSHITA DENKI SANGYO KK; (MATU) MATSUSHITA ELECTRIC IND CO  
LTD

CYC 4  
PI US 2004023115 A1 20040205 (200415)\* 8 H01M004-58  
JP 2004127913 A 20040422 (200428) 29 H01M004-02  
CN 1481041 A 20040310 (200437) H01M004-58  
KR 2004012541 A 20040211 (200438) H01M004-60  
ADT US 2004023115 A1 US 2003-620687 20030717; JP 2004127913 A JP  
2003-160969 20030605; CN 1481041 A CN 2003-152250 20030730; KR  
2004012541 A KR 2003-52624 20030730

PRAI JP 2002-223862 20020731

IC ICM H01M004-02; H01M004-58; H01M004-60  
ICS H01M002-02; H01M004-62; H01M004-66; H01M010-36; H01M010-40

AB US2004023115 A UPAB: 20040302  
NOVELTY - A lithium rechargeable battery comprises positive and negative electrodes, and a non-aqueous electrolyte. The negative electrode comprises a copper core material and a negative electrode material mixture layer fixed on the core material. The negative electrode material mixture layer includes an active material having a mixture of artificial graphite particles and spherical graphite particles.

DETAILED DESCRIPTION - A lithium rechargeable battery comprises positive and negative electrodes (2, 3), and a non-aqueous electrolyte. The negative electrode comprises a copper core material and a negative electrode material mixture layer fixed on the core material. The negative electrode material mixture layer includes an active material having a mixture of artificial and spherical graphite particles. The artificial graphite particles are isotropic artificial graphite particles with graphite structure oriented at random. They have not more than 3.362 Angstrom inter planar spacing d002 between the (002) planes; not more than 1000 I002/I110 ratio of the peak intensity (I002) attributed to the (002) plane to the peak intensity (I110) attributed to the (110) plane, which are found from a diffraction pattern of the artificial graphite particles A molded into a pellet of 1.6 g/cm<sup>3</sup> in density; 0.85-0.95 mean circularity of the particles; 15-30  $\mu$ m particle diameter (D50) corresponding to a volume fraction of 50% and 0.2-0.5 D10/D90 ratio of a particle diameter (D10) corresponding to a volume fraction of 10% to a particle diameter (D90) corresponding to a volume fraction of 90%; not less than 1 g/cm<sup>3</sup> tap density after tapping of 900 times with a stroke length of 18 mm; and not more than m<sup>2</sup>/g specific surface area. The spherical graphite particles B have a mean circularity of the particles of 0.88-1; D50 corresponding to a volume fraction of 50% of 5-15  $\mu$ m; d002 between the (002) planes of not more than 3.357 Angstrom; and not more than 8 m<sup>2</sup>/g specific surface area.

USE - As driving power source.

ADVANTAGE - The inventive battery is a portable, wireless, compact and lightweight battery having high energy density.

DESCRIPTION OF DRAWING(S) - The figure is an oblique view illustrating a lithium rechargeable battery.

Electrode group 1

Positive and negative electrodes 2, 3

Case 4

Sealing plate 5

Dwg.1/1

TECH US 2004023115 A1UPTX: 20040302

TECHNOLOGY FOCUS - CERAMICS AND GLASS - Preferred Material: The

artificial graphite particles A are obtained by kneading and granulating a base material of pulverized bulk mesophase pitch with pitch in a softened state and/or thermosetting resin, carbonizing the resulting granules at 700-1500degreesC, and graphitizing the carbonized granules at 2500-3000degreesC. The spherical graphite particles B are natural graphite particles and/or natural graphite particles partially subjected to reforming treatment to obtain amorphous surfaces. Preferred Composition: The spherical graphite particles B are present at 5-45 wt.%. Preferred Property: The negative electrode material mixture layer has a density of 1.6-1.8 g/cm<sup>3</sup> and a thickness of 40-100  $\mu$ m.

TECHNOLOGY FOCUS - ELECTRICAL POWER AND ENERGY - Preferred Component: The positive and negative electrodes with a separator are wound to form an electrode group (1). The electrode group is accommodated and sealed in a prismatic metal case (4) or a case made of aluminum foil laminate and a resin film.

TECHNOLOGY FOCUS - POLYMERS - Preferred Component: The negative electrode material mixture layer further includes a rubber-like binder containing a butadiene unit and a cellulose-based thickener. Preferred Composition: The rubber-like binder is added at not more than 3 pbw with respect to 100 pbw active material.

FS CPI EPI

FA AB; GI

MC CPI: A12-E06A; A12-E06C; L03-E01B3  
EPI: X16-E01C; X16-E08A

L30 ANSWER 4 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN

AN 2002-734690 [80] WPIX

DNN N2002-579196 DNC C2002-208090

TI Lead-acid battery for electric cars and elevators, comprises anode which is added with carbon and simple substance and/or compound having catalyst for desulfurization or oxidation of sulfur oxides.

DC E36 J04 L03 X16

IN HONBO, K; HOSHI, E; MURANAKA, Y; TAKEUCHI, S

PA (HITA) HITACHI LTD; (KOBE) SHIN KOBE ELECTRIC MACHINERY; (HONB-I) HONBO K; (HOSH-I) HOSHI E; (MURA-I) MURANAKA Y; (TAKE-I) TAKEUCHI S

CYC 28

PI EP 1248307 A1 20021009 (200280)\* EN 31 H01M004-14  
R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK  
NL PT RO SE SI TR

JP 2002367613 A 20021220 (200313) 15 H01M004-62

US 2003049528 A1 20030313 (200321) H01M004-62

US 2004180264 A1 20040916 (200461) H01M004-62

ADT EP 1248307 A1 EP 2002-5531 20020311; JP 2002367613 A JP 2002-67800  
20020313; US 2003049528 A1 US 2002-96505 20020313; US 2004180264 A1  
Cont of US 2002-96505 20020313, US 2004-812005 20040330

PRAI JP 2001-104080 20010403

IC ICM H01M004-14; H01M004-62

ICS B01J021-18; B01J023-00; H01M004-38

AB EP 1248307 A UPAB: 20021212

NOVELTY - The lead-acid battery comprises an anode, a cathode and an electrolytic solution. The anode is added with carbon, and a simple substance and/or compound having a catalyst for desulfurization or oxidation of sulfur oxides.

DETAILED DESCRIPTION - An INDEPENDENT CLAIM is included for the carbon material used in lead-acid batteries.

USE - For electrical cars, simple hybrid cars, power storage systems, elevators, electric tools, uninterruptable power sources

and distributed power sources.

ADVANTAGE - The lead-acid battery is superior in high-efficiency charging characteristics, due to the addition of carbon and simple substance and/or compound having a catalyst, to the anode. The conductivity of lead sulfate, and the solubility of lead sulfate into the lead ion are improved, hence the charging reaction of anode active material proceeds smoothly. The carbon material used in anode has superior charge acceptability. The battery exhibits a small energy loss due to the gas generated during large-current charging. The current value necessary to discharge the total discharge capacity of the battery in 0.5 hour and 1 hour, are 2C and 1C, respectively. The carbon can be adsorbed on the reaction interface of the active material of the lead-acid battery, thereby the passivation of lead sulfate which is called sulfation can be suppressed, no passivation proceeds even when complete discharge has been made and charge acceptability is remarkably improved.

DESCRIPTION OF DRAWING(S) - The figure shows the model of the catalyst used for desulfurization.  
Dwg. 6/8

TECH EP 1248307 A1 UPTX: 20021212  
TECHNOLOGY FOCUS - INORGANIC CHEMISTRY - Preferred Catalyst: The catalyst, is catalyst used for petroleum refining, fuel oil desulfurization, gas production or pollution control.  
Preferred Substance: The simple substance is an oxide, sulfate, hydroxide and/or carbide of hafnium, niobium, tantalum, tungsten, silver, zinc, nickel, copper, molybdenum, copper, vanadium, manganese, barium, potassium, cesium, rubidium, strontium and sodium. The carbon is carbon black, acetylene black, natural graphite, artificial graphite, pyrolytic carbon, coke, isotropic graphite, mesophase carbon, pitch-based carbon fiber, carbon fiber by vapor phase growth, carbon fluoride, nanocarbon, active carbon, active carbon fiber or polyacrylonitrile-based fiber.

FS CPI EPI  
FA AB; GI; DCN  
MC CPI: E11-Q02; E31-F01A; E31-N04; E31-N05A; E33; E34; E35; J04-E04;  
L03-E01B; L03-E01B1; N06  
EPI: X16-E04

L30 ANSWER 5 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN  
AN 2002-398958 [43] WPIX  
DNN N2002-312950 DNC C2002-112497  
TI Carbon film manufacture for electrodes of non-aqueous electrolyte secondary battery, involves dispersing carbon material in aprotic solvent in which electroconductive substrate and counter electrode are soaked.

DC L03 X12 X16  
PA (KOUR-I) KOURA N; (SHAF) SHARP KK  
CYC 1  
PI JP 2002063894 A 20020228 (200243)\* 16 H01M004-04  
ADT JP 2002063894 A JP 2000-251100 20000822  
PRAI JP 2000-251100 20000822  
IC ICM H01M004-04  
ICS C25B007-00; H01B001-04; H01M004-02; H01M004-58; H01M004-66;  
H01M010-40

AB JP2002063894 A UPAB: 20020709  
NOVELTY - A carbon material (2) is dispersed in an aprotic solvent (1) in which iodine is dissolved. An electroconductive substrate and a counter electrode are soaked in the solvent as cathode (31) and anode (32), respectively. A carbon film is made to deposit on the surface of electroconductive substrate by electrophoresis electrodeposition, when a direct flow electric field

is impressed between the **electrodes**.

DETAILED DESCRIPTION - An INDEPENDENT CLAIM is also included for non-aqueous electrolyte secondary battery.

USE - For **electrodes** of non-aqueous electrolyte secondary battery (claimed).

ADVANTAGE - The carbon film is directly deposited on the substrate by simple method. The carbon film has reduced weight and high industrial utility. The active material density of **electrode** formed from carbon film is increased.

DESCRIPTION OF DRAWING(S) - The figure shows outline view of manufacture of carbon film.

Aprotic solvent 1

Carbon material 2

Cathode 31

Anode 32

Dwg.1/8

TECH JP 2002063894 AUPTX: 20020709

TECHNOLOGY FOCUS - ORGANIC CHEMISTRY - Preferred Compounds: The aprotic solvent is acetone or acetyl acetone. The carbon material is chosen from artificial **graphite**, natural **graphite**, high crystalline **graphite**, **mesophase**

**pitch**, isotropic **pitch**, mesocarbon micro beads and coke. The electroconductive substrate comprises **copper**, nickel, stainless steel, brass, molybdenum and tungsten.

FS CPI EPI

FA AB; GI

MC CPI: L03-E01B9

EPI: X12-D01C; X16-B01F; X16-E01; X16-E01C; X16-E01G; X16-E02

L30 ANSWER 6 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN

AN 2000-507805 [46] WPIX

DNN N2000-375557 DNC C2000-152320

TI Negative **electrode** for non-aqueous electrolyte secondary battery, has carbon material with aggregated **graphite** powder and **mesophase pitch** type **graphite** fibers.

DC A14 A85 L03 X16

PA (KOBE) SHIN KOBE ELECTRIC MACHINERY

CYC 1

PI JP 2000156226 A 20000606 (200046)\* 5 H01M004-58

ADT JP 2000156226 A JP 1998-327923 19981118

PRAI JP 1998-327923 19981118

IC ICM H01M004-58

ICS H01M004-02; H01M004-62; H01M010-40

AB JP2000156226 A UPAB: 20000921

NOVELTY - The **electrode** consists of a rolled **copper** foil on which a mixture of carbon material and polyvinylidene fluoride resin binder is adhered. The carbon material consists of aggregated **graphite** powder and **mesophase pitch** type **graphite** fiber material.

USE - For non-aqueous secondary battery e.g. nickel-cadmium battery, lead battery, nickel hydrogen battery used in portable telephone, notebook personal computer.

ADVANTAGE - The mixture density and mixture adhesion are raised by the usage of aggregated **graphite** powder and **mesophase graphite** powder, thus irreversible capacitance and the gas evolution at high temperature is suppressed. The peeling and the crack of a mixture layer is prevented, thus excellent cycle property is obtained.

Dwg.0/0

FS CPI EPI

FA AB  
MC CPI: A04-E10B; A12-E06; L03-E01B3  
EPI: X16-B01F; X16-E01C; X16-E09

L30 ANSWER 7 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN  
AN 1989-238387 [33] WPIX  
DNN N1989-181475 DNC C1989-106466  
TI Electrochemical detector for high speed liq. chromatography - has  
cell with working **electrode** composed of synthetic polymer  
cpd. and graphite, counter-**electrode**, reference  
**electrode**, etc..  
DC A89 B04 J04 S03  
PA (ACOM-N) ACOM KK; (SEKI) SEKISUI CHEM IND CO LTD  
CYC 1  
PI JP 01173861 A 19890710 (198933)\* 7  
ADT JP 01173861 A JP 1987-335803 19871228  
PRAI JP 1987-335803 19871228  
IC G01N027-46; G01N030-64  
AB JP 01173861 A UPAB: 19930923  
An electrochemical detector for high speed liq chromatography having  
a cell with a working **electrode**, a counter  
**electrode** and a reference **electrode**, and detection  
part is claimed where the working **electrode** is constructed  
of a compn mainly consisting of a synthesised polymer cpd and  
**spherical graphite** carbon.  
In the compn, the amnt of the **spherical**  
**graphite** carbon is pref 20 - 80 wt %. The synthesised  
polymer cpd is not specially limited, but it must be a material  
which is not dissolved or swelled with the eluate of the liq  
chromatography. For example, fluorine resin, polyphenylene sulphide  
resin etc are pref.  
USE/ADVANTAGE - An electrochemical detector which enables the  
measurement of high sensitivity is obtd. It is esp useful for  
measuring vitamins, steroids, and various chemicals in a sample  
which is sepd by high speed liq chromatography.  
0/0

FS CPI EPI  
FA AB; DCN  
MC CPI: A12-E14; A12-L04; A12-V03C2; B03-L; B04-B02D; B04-C03; B05-C06;  
B11-C08B; B11-C08D2; B12-K04; J04-B01C  
EPI: S03-E03; S03-E03C; S03-E09C5

L30 ANSWER 8 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN  
AN 1989-123644 [17] WPIX  
DNN N1989-094269 DNC C1989-054780  
TI Exothermic conductive coating compsns. - contg. spherical carbon  
particles, and synthetic resin, used in heating units with uniform  
temp. distribution up to 450 deg. C.  
DC A28 A32 G02 L03 P73 X25 X27  
IN OTA, T  
PA (ASKH-N) ASK HEATING KK; (OTAT-I) OTA T; (ASKH-N) ASK HEATING CO  
LTD; (OHOT-I) OHOTA T  
CYC 12  
PI EP 312688 A 19890426 (198917)\* EN 17  
R: CH DE FR GB LI NL SE  
JP 01107488 A 19890425 (198922)  
JP 01108276 A 19890425 (198923)  
NO 8802116 A 19890516 (198925)  
EP 312688 B1 19930107 (199302) EN 18 C09D005-24  
R: CH DE FR GB LI NL SE  
DE 3877314 G 19930218 (199308) C09D005-24  
CA 1323717 C 19931026 (199349) C09D005-24

NO 173878 B 19931108 (199350) C09D005-24  
 JP 06089270 B2 19941109 (199443) 7 C09D005-24  
 KR 9401521 B1 19940223 (199503) C09D005-24  
 US 5407741 A 19950418 (199521) 14 B32B005-16  
 ADT EP 312688 A EP 1988-107673 19880513; JP 01107488 A JP 1987-263955  
 19871021; JP 01108276 A JP 1987-263954 19871021; EP 312688 B1 EP  
 1988-107673 19880513; DE 3877314 G DE 1988-3877314 19880513, EP  
 1988-107673 19880513; CA 1323717 C CA 1988-566767 19880513; NO  
 173878 B NO 1988-2116 19880513; JP 06089270 B2 JP 1987-263954  
 19871021; KR 9401521 B1 KR 1988-5615 19880513; US 5407741 A Cont of  
 US 1988-192143 19880510, Cont of US 1991-663086 19910228, US  
 1993-13177 19930129  
 FDT DE 3877314 G Based on EP 312688; NO 173878 B Previous Publ. NO  
 8802116; JP 06089270 B2 Based on JP 01108276  
 PRAI JP 1987-263955 19871021; JP 1987-263954 19871021  
 REP 4.Jnl.Ref; A3...199101; JP 52115445; JP 60008377; JP 61235471;  
 No-SR.Pub; US 4035265  
 IC ICM B32B005-16; C09D005-24  
 AB ICS C09D005-00; H01B001-24; H05B003-14  
 EP 312688 A UPAB: 20010910  
 An exothermic conductive coating compsn comprises (i) carbon  
 particles which mainly (pref at least 60 wt%) consist of spherical  
 particles of dia not more than 500 microns; and (ii) a synthetic  
 resin, pref a polyester resin, epoxy resin, polyamide, polyimide,  
 polyethylene, F-contg polymer, polyetheretherketone, poly-phenylene  
 sulphide, silicone resin or poly-titanocarbosilane resin and pref as  
 25-220 pts wt per 100 pts wt of (a).  
 Pref particles (i) have been heat treated at least 1500 deg.C  
 and have an interplanar spacing in the crystalline structure of  
 3.425-3.358 Angstroms.  
 USE - Also claimed are conductive heating units which comprise  
 a film of (i) plus (ii) on a shaped solid surface having  
 electrode terminals mounted on it; in a claimed variant  
 there may be further laminated exothermic layers, each of which has  
 electrode terminals. The heating units can be used in  
 interior walls, flooring, roofing, furnace inner surfaces, pipe  
 surfaces, carpets, blankets etc.  
 ADVANTAGE - The compsns show uniform temp distribution and can  
 produce arbitrarily adjustable temps up to 450 deg.C. Heating units  
 can be prepd from complex-shaped substrates (e.g. with holes or  
 surface unevenness) without localised over heating (cf use of  
 needle-like, flake or fibrous carbon particles which does cause  
 overheating).  
 Dwg.12/12  
 FS CPI EPI GMPI  
 FA AB; GI  
 MC CPI: A12-B01; A12-E10; G02-A05; G02-A05B; L03-A02E; L03-H04A  
 EPI: X25-B01B; X27-E01A3; X27-E02  
 L30 ANSWER 9 OF 9 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN  
 AN 1975-H6109W [30] WPIX  
 TI Electric or solar energy heat store - has graphite and coal storage  
 medium contained under vacuum in reflector-walled vessel.  
 DC Q55  
 PA (SCHM-I) SCHMID A W  
 CYC 1  
 PI DE 2401859 A 19750717 (197530)\*  
 PRAI DE 1974-2401859 19740116  
 IC F03G007-02  
 AB DE 2401859 A UPAB: 19930831  
 The store has a central graphite electrode surrounded by  
 the storage medium in the form of spherical elements of

graphite or synthetic coal. The medium is contained in a graphite container surrounded by heat reflectors made of relatively thin graphite cylinders. The assembly is contained in an evacuated steel cylinder which also encloses a heat exchanger through which heat can be withdrawn from the store. In a modification the sun is used as a heat source, the rays being entered into the store through mirrors. The use of graphite or coal as the storage medium allows storage temperatures up to 3273 deg. K.

FS GMPI  
FA AB

=> file hcaplus

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FILE LAST UPDATED: 20 Dec 2005 (20051220/ED)

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L5	191171	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	GRAPHITE# OR BLACK (2A) LEAD OR PLUMBAGO## OR L2
L6	2533705	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	ARTIFICIAL? OR SYNTHETIC ? OR SIMULAT? OR FABRICAT? OR MANMADE OR MAN (A) MADE OR ERSATZ OR SUBSTITUT?
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L12	133	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	L7 (L) L5 AND L6 (L) L5 (L) L4
L13	18855	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	MESOPHASE OR MESOPHASIC
L14	48953	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	PITCH##
L16	2	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	L13 AND L12
L19	6	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	L13 AND L14 AND L4 AND L5 AND L3
L20	51	SEA FILE=HCAPLUS	ABB=ON	PLU=ON	L13 (L) L14 (L) L4 (L) L5
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        25 SEA FILE=HCAPLUS ABB=ON PLU=ON L22 AND (57? OR 51?)/SC

L26      25 SEA FILE=HCAPLUS ABB=ON PLU=ON L23 NOT (L9 OR L16 OR
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        OR L28

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=> file japio

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 L23 25 SEA FILE=HCAPLUS ABB=ON PLU=ON L22 AND (57? OR 51?)/SC  
  
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 L13 18855 SEA FILE=HCAPLUS ABB=ON PLU=ON MESOPHASE OR MESOPHASIC  
  
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 L16 2 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 AND L12  
 L19 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 AND L14 AND L4 AND  
 L5 AND L3  
 L20 51 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 (L) L14 (L) L4 (L)  
 L5  
 L21 24 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 (L) L14 (L) L4 (L)  
 L5 AND ELECTRO?/SC,SC  
 L22 27 SEA FILE=HCAPLUS ABB=ON PLU=ON L20 NOT L21  
 L23 25 SEA FILE=HCAPLUS ABB=ON PLU=ON L22 AND (57? OR 51?)/SC  
  
 L26 25 SEA FILE=HCAPLUS ABB=ON PLU=ON L23 NOT (L9 OR L16 OR  
 L19 OR L21)  
 L28 10 SEA FILE=HCAPLUS ABB=ON PLU=ON L26 AND (ELECTRODE# OR  
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 L32 0 SEA FILE=JICST-EPLUS ABB=ON PLU=ON L9 OR L16 OR L19 OR  
 L21 OR L28

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 PROCESSING COMPLETED FOR L31  
 PROCESSING COMPLETED FOR L33  
 PROCESSING COMPLETED FOR L34  
 L35 53 DUP REM L29 L31 L33 L34 (3 DUPLICATES REMOVED)

=> d all 135 1-53

L35 ANSWER 1 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2005:523824 HCAPLUS  
 DN 143:46084  
 ED Entered STN: 17 Jun 2005  
 TI A process of production of carbon negative electrode  
 material for lithium ion battery  
 IN Ma, Junqi; Wang, Jianqiao; Fu, Zhenming; Ding, Xiaoyang; Li,  
 Shaoming; Zhang, Dianhao; Feng, Suning  
 PA Shanghai Shanshan Tech Co., Ltd., Peop. Rep. China  
 SO PCT Int. Appl., 13 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA Chinese  
 IC ICM H01M004-58  
 ICS H01M004-04; C01B031-00  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 49, 51

FAN.CNT 1		PATENT NO.		KIND	DATE	APPLICATION NO.	DATE
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PI	WO 2005055346		A1	20050616	WO 2004-CN1297		20041115

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA,  
 CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,  
 GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP,  
 KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,  
 MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD,  
 SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,  
 VC, VN, YU, ZA, ZM, ZW  
 RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW,  
 AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ,

DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL,  
 PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN,  
 GQ, GW, ML, MR, NE, SN, TD, TG

PRAI CN 2003-10108982 A 20031201

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
WO 2005055346	ICM	H01M004-58
	ICS	H01M004-04; C01B031-00
	IPCI	H01M0004-58 [ICM,7]; H01M0004-04 [ICS,7]; C01B0031-00 [ICS,7]

AB The present invention provides a carbon neg. **electrode** material for lithium ion battery by using natural **graphite**, esp. **spherical natural graphite** as raw material A, coal tar/coal pitch/petroleum pitch or mixt. thereof as raw material B, mixing the raw materials in a ratio of 1/0.5-10 of A to B, the resultant mixt. subjecting to polymn. reaction at a temp. of 350-500 °C under the pressure of 0.01 to 10 MPa for 5 to 420 min, drying the resultant product after extn. and sepn., the dried product having the wt. gain of 2-100 % compared with raw material of graphite, finally, carbonizing or graphitizing the product to obtain carbon neg. **electrode** material having a coated layer of artificial carbon or **artificial graphite**. The carbon neg. **electrode** material for lithium ion battery has advantages of high specific capacity, long cycle lifetime and the like, is a desirable carbon neg. **electrode** material.

ST carbonization graphitization graphite coal tar petroleum pitch battery anode

IT Electric capacitance  
(high specific capacity; process of prodn. of carbon neg. **electrode** material for lithium ion battery)

IT Secondary batteries  
(lithium; process of prodn. of carbon neg. **electrode** material for lithium ion battery)

IT Polymerization  
(of graphite and pitches; process of prodn. of carbon neg. **electrode** material for lithium ion battery)

IT Battery anodes  
Carbonization  
Coal tar pitch  
Drying  
Extraction  
Graphitization  
Petroleum pitch  
(process of prodn. of carbon neg. **electrode** material for lithium ion battery)

IT Coal tar  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(process of prodn. of carbon neg. **electrode** material for lithium ion battery)

IT 7782-42-5P, Graphite, uses  
RL: DEV (Device component use); IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)  
(process of prodn. of carbon neg. **electrode** material for lithium ion battery)

IT 7440-44-0, Carbon, uses  
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
(process of prodn. of carbon neg. **electrode** material

for lithium ion battery)

RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

- (1) Fudan University; CN 1274957 A 2000 HCAPLUS
- (2) He, M; Carbon 2002, V3, P3
- (3) Jiang, H; Carbon 2001, V3, P12
- (4) Shanghai Shanshan Tech Co Ltd; CN 1382624 A 2002 HCAPLUS
- (5) Song, W; Chinese J Power Sources 2002, V26(6), P428 HCAPLUS
- (6) Song, W; Study on composite materials prepared by coal pitch coating on natural flake Graphite used as carbon cathode materials for lithium ion secondary batteries 2001, V20(6), P27

L35 ANSWER 2 OF 53 INSPEC (C) 2005 IEE on STN

AN 2005:8473336 INSPEC DN A2005-16-8245-003; B2005-08-8410E-015

TI Study of **mesophase pitch-based graphite**

foam used as anodic materials in lithium ion rechargeable batteries.

AU Jie Yang; Zeng-Min Shen; Rui-Sheng Xue (Inst. of Carbon Fibers & Composites, Beijing Univ. of Chem. Technol., China); Zhi-Biao Hao

SO Journal of Materials Science (1 March 2005) vol.40, no.5, p.1285-7.

15 refs.

Published by: Kluwer Academic Publishers

CODEN: JMTSAS ISSN: 0022-2461

SICI: 0022-2461(20050301)40:5L:1285:SMPB;1-5

DT Journal

TC Experimental

CY United States

LA English

AB We attempt to investigate the **mesophase pitch** carbon foams heat treated at high temperature as anode materials for lithium ion secondary batteries. The foaming technique used in this study is similar to that reported by Klett et al where powders of **mesophase pitch** were synthesized from FCC decant oil. The foams were characterized by X-ray diffraction. The **electrodes** were prepared by coating slurries of the carbon foams powder (90 wt%) polyvinylidene fluoride (PVDF, 10 wt%) dissolved in N-methyl pyrrolidinone (NMP) on copper foil. The carbon foams exhibit a superior anode performance with stable capacitance that can reach as high as 306 mAhg<sup>-1</sup>.

CC A8245 Electrochemistry and electrophoresis; A8630F Secondary cells;

A8270R Aerosols and foams; A8120V Preparation of fullerenes and

fullerene-related materials, intercalation compounds, and diamond;

A8270K Emulsions and suspensions; A8140G Other heat and

thermomechanical treatments; B8410E Secondary cells

CT CAPACITANCE; ELECTROCHEMICAL **ELECTRODES**; FOAMS;

**GRAPHITE**; HEAT TREATMENT; SECONDARY CELLS; SLURRIES; X-RAY

DIFFRACTION

ST **mesophase pitch-based graphite foam**; anodic materials; lithium ion rechargeable batteries; heat treatment; lithium ion secondary batteries; foaming technique; powders; FCC decant oil; X-ray diffraction; slurries; polyvinylidene fluoride; N-methyl pyrrolidinone; capacitance; C

CHI C el

ET N; C

L35 ANSWER 3 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:100617 HCAPLUS

DN 140:114301

ED Entered STN: 08 Feb 2004

TI Lithium rechargeable battery

IN Kato, Fumio; Oura, Takafumi; Fukumoto, Yusuke; Yamamoto, Norihiro;

Fujiwara, Shozo

PA Matsushita Electric Industrial Co., Ltd., Japan

SO U.S. Pat. Appl. Publ., 18 pp.

CODEN: USXXCO

DT Patent

LA English

IC ICM H01M004-58

ICS H01M004-62; H01M004-66

INCL 429231800; 429245000; 429217000

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004023115	A1	20040205	US 2003-620687	20030717
JP 2004127913	A2	20040422	JP 2003-160969	20030605
CN 1481041	A	20040310	CN 2003-152250	20030730

PRAI JP 2002-223862 A 20020731

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
US 2004023115	ICM	H01M004-58
	ICS	H01M004-62; H01M004-66
	INCL	429231800; 429245000; 429217000
	IPCI	H01M0004-58 [ICM,7]; H01M0004-62 [ICS,7]; H01M0004-66 [ICS,7]
JP 2004127913	NCL	429/231.800
	IPCI	H01M0004-02 [ICM,7]; H01M0002-02 [ICS,7]; H01M0004-58 [ICS,7]; H01M0004-62 [ICS,7]; H01M0004-66 [ICS,7]; H01M0010-40 [ICS,7]
	FTERM	5H011/AA03; 5H011/AA09; 5H011/AA13; 5H011/BB03; 5H011/CC06; 5H011/CC10; 5H017/AA03; 5H017/AS01; 5H017/AS10; 5H017/CC01; 5H017/EE01; 5H017/HH05; 5H029/AJ02; 5H029/AJ03; 5H029/AJ05; 5H029/AJ12; 5H029/AK03; 5H029/AL07; 5H029/AM03; 5H029/AM05; 5H029/AM07; 5H029/BJ02; 5H029/BJ14; 5H029/CJ02; 5H029/CJ08; 5H029/DJ02; 5H029/DJ08; 5H029/DJ16; 5H029/DJ17; 5H029/EJ01; 5H029/EJ03; 5H029/EJ13; 5H029/HJ01; 5H029/HJ05; 5H029/HJ07; 5H029/HJ08; 5H029/HJ13; 5H029/HJ14; 5H050/AA02; 5H050/AA06; 5H050/AA07; 5H050/AA08; 5H050/AA15; 5H050/AA19; 5H050/BA17; 5H050/CA08; 5H050/CB08; 5H050/DA11; 5H050/EA28; 5H050/FA05; 5H050/FA17; 5H050/FA19; 5H050/GA02; 5H050/GA05; 5H050/GA06; 5H050/GA10; 5H050/HA01; 5H050/HA05; 5H050/HA07; 5H050/HA08; 5H050/HA13; 5H050/HA14
CN 1481041	IPCI	H01M0004-58 [ICM,7]; H01M0004-62 [ICS,7]; H01M0010-36 [ICS,7]; H01M0010-40 [ICS,7]

AB The present invention provides a lithium rechargeable battery including a neg. electrode comprising a copper core material, to which is fixed an active material made by mixing artificial graphite particles obtained by kneading and granulating a base material of pulverized bulk mesophase pitch with pitch in a softened state and/or thermosetting resin, carbonizing the resulting granules at 700 to 1500° and graphitizing the carbonized granules at 2500 to 3000° with spherical graphite particles

having a high mean circularity, with a view to inhibiting the deterioration in battery capacity through the repeated charge/discharge cycles, which occurs remarkably in a high energy d. lithium rechargeable battery, and providing excellent discharge characteristic and safety.

ST battery lithium secondary; safety lithium rechargeable battery  
 IT Battery anodes  
 Battery cathodes  
 Coal tar pitch  
 Mesophase pitch  
 (lithium rechargeable battery)

IT Coal tar  
 RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
 (lithium rechargeable battery)

IT Secondary batteries  
 (lithium; lithium rechargeable battery)

IT Plastics, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (thermosetting; lithium rechargeable battery)

IT 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate  
 623-53-0, Ethyl methyl carbonate 7440-50-8, Copper  
 , uses 7782-42-5, Graphite, uses 12190-79-3,  
 Cobalt lithium oxide colio2 21324-40-3, Lithium  
 hexafluorophosphate  
 RL: DEV (Device component use); USES (Uses)  
 (lithium rechargeable battery)

L35 ANSWER 4 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2004:900943 HCAPLUS  
 DN 141:382112  
 ED Entered STN: 28 Oct 2004  
 TI Production of spherical graphite particles and its use in lithium  
 ion secondary battery cathode material  
 IN Sakai, Minoru; Nagayama, Katsuhiko; Hatano, Hitomi; Morioka,  
 Hironori; Honma, Makoto  
 PA JFE Chemical Corporation, Japan  
 SO Jpn. Kokai Tokkyo Koho, 17 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM C01B031-04  
 ICS H01M004-02; H01M004-58; H01M010-40  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
 Technology)  
 Section cross-reference(s): 49

FAN.CNT 1	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2004299944	A2	20041028	JP 2003-93308	200303 31

PRAI JP 2003-93308

20030331

# CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2004299944	ICM	C01B031-04
	ICS	H01M004-02; H01M004-58; H01M010-40
	IPCI	C01B0031-04 [ICM,7]; H01M0004-02 [ICS,7]; H01M0004-58 [ICS,7]; H01M0010-40 [ICS,7]
	FTERM	4G146/AA02; 4G146/AA23; 4G146/AB01; 4G146/AB02;

4G146/AC02A; 4G146/AC02B; 4G146/AC17A;  
 4G146/AC17B; 4G146/AC22A; 4G146/AC22B;  
 4G146/AD03; 4G146/AD11; 4G146/AD25; 4G146/BA22;  
 4G146/BA23; 4G146/BA43; 4G146/BA49; 4G146/BB03;  
 4G146/BB06; 4G146/BB18; 4G146/BC04; 4G146/BC32A;  
 4G146/BC32B; 5H029/AJ03; 5H029/AJ05; 5H029/AK02;  
 5H029/AK03; 5H029/AK05; 5H029/AK08; 5H029/AL07;  
 5H029/AM02; 5H029/AM03; 5H029/AM04; 5H029/AM05;  
 5H029/AM07; 5H029/AM16; 5H029/BJ03; 5H029/CJ02;  
 5H029/CJ08; 5H029/EJ12; 5H029/HJ01; 5H029/HJ05;  
 5H029/HJ06; 5H029/HJ08; 5H029/HJ10; 5H029/HJ13;  
 5H029/HJ14; 5H050/AA07; 5H050/AA08; 5H050/BA17;  
 5H050/CA02; 5H050/CA11; 5H050/CA16; 5H050/CB08;  
 5H050/EA24; 5H050/GA02; 5H050/GA10; 5H050/HA01;  
 5H050/HA05; 5H050/HA06; 5H050/HA08; 5H050/HA13;  
 5H050/HA14

- AB Spherical **graphite** particles are produced from **mesophase pitch** micro beads (av. grain diam. 20-60  $\mu\text{m}$ ) by calcining at 350-450° to obtain **graphite** precursors contg. 4-20 wt.% volatile matter, and then graphitizing at  $\geq 2000^\circ$ , preferably 2800-3200° to obtain **cryst. graphite** particles having apparent sp. gr. 6.0 (detd. by JIS R7222-1997), d. 2.210-2.240 g/cm<sup>3</sup>, and inner pore diam. 0.1-30  $\mu\text{m}$ . The secondary battery **cathode** has larger service capacity around cubic measure, and does not cause deterioration in cycle quality.
- ST secondary lithium battery cathode graphite particle size control
- IT **Mesophase pitch**  
 (coal tar; prodn. of spherical **graphite** particles and its use in lithium ion secondary battery **cathode** material)
- IT Coal tar **pitch**  
 (**mesophase**; prodn. of spherical **graphite** particles and its use in lithium ion secondary battery **cathode** material)
- IT Battery cathodes  
 (prodn. of spherical graphite particles and its use in lithium ion secondary battery cathode material)
- IT 7440-44-0, Carbon, uses  
 RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)  
 (meso phase; micro beads; prodn. of spherical graphite particles and its use in lithium ion secondary battery cathode material)
- IT 7782-42-5P, Graphite, preparation  
 RL: IMF (Industrial manufacture); PREP (Preparation)  
 (prodn. of spherical graphite particles and its use in lithium ion secondary battery cathode material)
- L35 ANSWER 5 OF 53 INSPEC (C) 2005 IEE on STN DUPLICATE 1
- AN 2005:8414410 INSPEC DN A2005-13-8245-011; B2005-07-8410E-001
- TI Anomalous improvement of the electrochemical properties of mesocarbon microbeads by Ar-H<sub>2</sub>-SF<sub>6</sub> thermal plasma treatment.
- AU Tanaka, H.; Xu, J.Y. (Nat. Inst. for Mater. Sci., Ibaraki, Japan); Kurihara, M.; Maruyama, S.; Ohashi, N.; Moriyoshi, Y.; Ishigaki, T.
- SO Carbon (2004) vol.42, no.15, p.3229-35. 22 refs.  
 Published by: Elsevier  
 Price: CCCC 0008-6223/2004/\$30.00  
 CODEN: CRBNAH ISSN: 0008-6223  
 SIC1: 0008-6223(2004)42:15L.3229:AI EP;1-5
- DT Journal
- TC Practical; Experimental



CY United Kingdom  
 LA English  
 AB Highly graphitized mesocarbon microbead (MCMB) powders, which consist of **spherical artificial graphite** particles, were treated in Ar-H<sub>2</sub>-SF<sub>6</sub> radio-frequency inductively coupled thermal plasma. The results of X-ray diffractometry, scanning electron microscopy, Fourier transform infrared spectroscopy, Raman spectroscopy, X-ray photoelectron spectroscopy, and element analysis showed that the plasma treatment greatly affected the surface structure and chemical composition of the MCMB powders. Sulfur and fluorine could be incorporated into the particle by treating them with plasma containing SF<sub>6</sub>, and this plasma modification improved the electrochemical properties of the powders. A lithium-ion rechargeable cell with a negative **electrode** of plasma-treated powder had a discharge capacity as much as 13% greater than that of a cell with a negative **electrode** of untreated powder, while its irreversible capacity loss (ICL) was as much as 9% less. The specific ICL calculated by dividing the ICL by surface area decreased as the BET surface area increased. Furthermore, a new plateau near 2 V was added to the discharge curve by treating the MCMB powder with SF<sub>6</sub> plasma but not by treating it with Ar-H<sub>2</sub>, Ar-N<sub>2</sub>, Ar-H<sub>2</sub>-CO<sub>2</sub>, or Ar-N<sub>2</sub>-CO<sub>2</sub> plasmas. The incorporation of sulfur and fluorine should be a key factor for the improvement of negative **electrode** characteristics.

CC A8245 Electrochemistry and electrophoresis; A8280P Electron spectroscopy for chemical analysis (photoelectron, Auger spectroscopy, etc.); A7830G Infrared and Raman spectra in inorganic crystals; A7960 Photoemission and photoelectron spectra (condensed matter); A5250G Plasma heating; A8630F Secondary cells; A6820 Solid surface structure; B8410E Secondary cells

CT ELECTROCHEMICAL **ELECTRODES**; ELECTROCHEMISTRY; FLUORINE; FOURIER TRANSFORM SPECTRA; GRAPHITE; INFRARED SPECTRA; PLASMA RADIOFREQUENCY HEATING; POWDERS; RAMAN SPECTRA; SCANNING ELECTRON MICROSCOPY; SECONDARY CELLS; SULPHUR; SURFACE STRUCTURE; X-RAY DIFFRACTION; X-RAY PHOTOELECTRON SPECTRA

ST electrochemical properties; mesocarbon microbeads; **spherical artificial graphite particles**; X-ray diffractometry; scanning electron microscopy; Fourier transform infrared spectroscopy; Raman spectroscopy; X-ray photoelectron spectroscopy; element analysis; surface structure; chemical composition; irreversible capacity loss; sulfur; fluorine; **negative electrode**; radio-frequency inductively coupled thermal plasma; Li; C

CHI Li el; C sur, C el  
 ET Ar\*F\*H\*S; SF<sub>6</sub>; S cp; cp; F cp; Ar-H<sub>2</sub>-SF<sub>6</sub>; F\*S; Ar\*H; Ar-H<sub>2</sub>; Ar\*N; Ar-N<sub>2</sub>; C\*H\*Ar\*O; CO<sub>2</sub>; C cp; O cp; Ar-H<sub>2</sub>-CO<sub>2</sub>; C\*Ar\*N\*O; Ar-N<sub>2</sub>-CO<sub>2</sub>; Li; C

L35 ANSWER 6 OF 53 COMPENDEX COPYRIGHT 2005 EEI on STN  
 AN 2004(31):7670 COMPENDEX  
 TI Effect of intercalation on electrical and mechanical properties of C/C composites.  
 AU Macherzynska, B. (Department of Advanced Ceramics Fac. of Mat. Science and Ceramics AGH Univ. of Science and Technology, 30-059 Crakow, Poland); Blazewicz, S.  
 SO Journal of Physics and Chemistry of Solids v 65 n 10 October 2004  
 2004.p 1745-1750  
 CODEN: JPCSAW ISSN: 0022-3697  
 PY 2004  
 DT Journal  
 TC Experimental  
 LA English  
 AB The work is concerned with modification of C/C composites by

intercalation of **copper** chloride. The samples of composites were made from **graphite** fibres and carbon matrix derived from **mesophase pitch** and from phenol-formaldehyde resin. The samples were prepared by impregnating **graphite** fibres with a liquid **pitch** or polymer solution to obtain unidirectional laminates. The laminates were used to prepare the composites which were then subjected to carbonization and graphitization up to 2150deg C. The work discusses the problem relevant to the effect of intercalation on mechanical and electrical properties of composites. The studies indicated that both **mesophase pitch**-based composites and phenolic-derived carbon-carbon composites changed their electrical and mechanical properties upon intercalation with **copper** chloride. Electrical conductivity of both types of composites decreased as a result of the damages formed during intercalation. \$CPY 2004 Elsevier Ltd. All rights reserved. 17 Refs.

CC 415.4 Other Structural Materials; 701.1 Electricity: Basic Concepts and Phenomena; 704.1 Electric Components; 801.4.1 Electrochemistry; 802.2 Chemical Reactions  
CT \*Carbon carbon composites; **Electrodes**; Electrochemistry; Mechanical properties; Graphitization; Electric conductivity; Carbonization; Intercalation compounds  
ST Gaseous precursors; Electrochemical devices; Acceptor molecules

L35 ANSWER 7 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:773819 HCAPLUS

DN 139:294631

ED Entered STN: 03 Oct 2003

TI Lithium ion secondary battery showing good charge-discharge cycle characteristics

IN Nishikawa, Reiji; Yoda, Kiyoto; Suzuki, Masami; Shikoda, Masataka

PA Toshiba Battery Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01M010-40

ICS H01M004-48; H01M004-62

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2003282147	A2	20031003	JP 2002-86994	20020326

PRAI JP 2002-86994

20020326

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2003282147	ICM	H01M010-40
	ICS	H01M004-48; H01M004-62
	IPCI	H01M0010-40 [ICM,7]; H01M0004-48 [ICS,7]; H01M0004-62 [ICS,7]

AB The title battery is equipped with: an anode which contains niobium pentaoxide, electroconductive materials, and a binder; **cathode** which carbon material electrochem. binding and releasing lithium and a binder; and nonaq. electrolyte, wherein the electroconductive material of the the anode is made of a mixt. of carbon black chosen from acetylene black, ketjen black, oil black, furnace black and **graphite** chosen from natural

graphite, artificial graphite, expanded  
 graphite, mesophase graphite  
 pitch, mesophase graphite fiber and  
 wherein the binder of cathode is made of a mixt. of rubber  
 polymer chosen from SBR, NBR, and MBR and cellulose or CMC or  
 polyacrylic acid deriv. chosen from polyacrylic acid, ammonium  
 polyacrylate, and lithium polyacrylate. The battery shows excellent  
 properties on the charge-discharge cycle.  
 lithium ion secondary battery  
 ST Anodes  
 IT Cathodes  
 (lithium ion secondary battery)  
 IT Carbon black, uses  
 Styrene-butadiene rubber, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (lithium ion secondary battery)  
 IT Secondary batteries  
 (lithium ion; lithium ion secondary battery)  
 IT Carbon fibers, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (mesophase pitch-based; lithium ion secondary battery)  
 IT 1313-96-8, Niobium pentaoxide 7782-42-5, Graphite, uses  
 9003-01-4, Polyacrylic acid  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (lithium ion secondary battery)  
 IT 9003-55-8  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (styrene-butadiene rubber, lithium ion secondary battery)

L35 ANSWER 8 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2002:964681 HCAPLUS  
 DN 138:42042  
 ED Entered STN: 20 Dec 2002  
 TI Improved cathodes for alkaline batteries  
 IN Zeng, Shuming  
 PA The Gillette Company, USA  
 SO PCT Int. Appl., 32 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 IC ICM H01M004-62  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
 Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002101856	A2	20021219	WO 2002-US17746	20020606
WO 2002101856	A3	20040401		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
US 2003008211	A1	20030109	US 2001-880651	

200106  
13

US 6596438 B2 20030722  
EP 1425812 A2 20040609 EP 2002-739693

200206  
06

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,  
PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

BR 2002010308 A 20040928 BR 2002-10308

200206  
06

JP 2005518633 T2 20050623 JP 2003-504491

200206  
06

PRAI US 2001-880651 A1 20010613  
WO 2002-US17746 W 20020606

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
WO 2002101856	ICM	H01M004-62
	IPCI	H01M0004-62 [ICM,7]
US 2003008211	IPCI	H01M0004-62 [ICM,7]; H01M0004-50 [ICS,7]; H01M0004-42 [ICS,7]; H01M0010-26 [ICS,7]
	NCL	429/229.000
EP 1425812	IPCI	H01M0004-62 [ICM,7]
BR 2002010308	IPCI	H01M0004-62 [ICM,7]
JP 2005518633	IPCI	H01M0006-08 [ICM,7]; H01M0004-50 [ICS,7]; H01M0004-62 [ICS,7]
	FTERM	5H024/AA03; 5H024/AA14; 5H024/BB01; 5H024/CC02; 5H024/DD17; 5H024/EE03; 5H024/FF08; 5H024/HH00; 5H024/HH01; 5H024/HH11; 5H024/HH13; 5H050/AA02; 5H050/AA12; 5H050/BA03; 5H050/CA05; 5H050/CB13; 5H050/DA10; 5H050/EA09; 5H050/FA07; 5H050/FA16; 5H050/GA02; 5H050/HA01; 5H050/HA04; 5H050/HA07; 5H050/HA13; 5H050/HA14
AB		The title battery has a cathode comprising manganese dioxide and graphitized carbon fibers with an av. BET surface area of 10-60 m <sup>2</sup> /g. Optionally, graphitized mesophase pitch-based carbon fibers can be used instead of graphitized carbon fibers. The graphitized mesophase pitch-based carbon fibers may be heat treated at 800-1200° with KOH. The use of graphitized carbon fibers in the cathode increases cathode cond. and results in improved battery performance.
ST		battery alk graphitized carbon fiber cathode; graphitized mesophase pitch based carbon fiber cathode battery
IT		Battery cathodes Primary batteries (cathodes contg. graphitized carbon fibers for alk. batteries)
IT		Carbon fibers, uses RL: DEV (Device component use); USES (Uses) (graphite, mesophase pitch-based, TA 1F, DKE-X, 42C, 42D, LB 3F; cathodes contg. graphitized carbon fibers for alk. batteries)
IT		Carbon fibers, uses RL: DEV (Device component use); USES (Uses) (graphite; cathodes contg. graphitized carbon fibers for alk. batteries)
IT		7782-42-5, Graphite, uses RL: DEV (Device component use); USES (Uses) (DCN 2, MP 0702X; cathodes contg. graphitized carbon fibers and flaky cryst. or expanded graphite for alk. batteries)
IT		7440-66-6, Zinc, uses

RL: DEV (Device component use); USES (Uses)  
 (anode; cathodes contg. graphitized carbon fibers for alk.  
 batteries with)  
 IT 1310-58-3, Potassium hydroxide (KOH), uses  
 RL: DEV (Device component use); NUU (Other use, unclassified); USES  
 (Uses)  
 (electrolyte and heat treating agent; cathodes contg. graphitized  
 carbon fibers for alk. batteries)  
 IT 1313-13-9, Manganese dioxide, uses  
 RL: DEV (Device component use); USES (Uses)  
 (electrolytic, cathode; cathodes contg. graphitized carbon fibers  
 for alk. batteries)

L35 ANSWER 9 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2002:172340 HCAPLUS  
 DN 136:202906  
 ED Entered STN: 08 Mar 2002  
 TI Processes for producing coke, artificial graphite and carbon  
 material for negative electrode of non-aqueous solvent  
 type secondary battery and pitch composition used therefor  
 IN Kanno, Koichi; Tsuruya, Hirotaka; Fujiura, Ryuji  
 PA Japan  
 SO U.S. Pat. Appl. Publ., 6 pp.  
 CODEN: USXXCO

DT Patent  
 LA English

IC ICM C10B047-00  
 ICS C10G001-00

INCL 201001000  
 CC 51-10 (Fossil Fuels, Derivatives, and Related Products)  
 Section cross-reference(s): 52, 57

FAN.CNT	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2002027066	A1	20020307	US 2001-923373	20010808
	JP 2002083595	A2	20020322	JP 2000-270315	20000906
	EP 1186646	A2	20020313	EP 2001-119188	20010809
	EP 1186646	A3	20030502		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
PRAI	JP 2000-270315	A	20000906		

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
US 20020027066	ICM	C10B047-00
	ICS	C10G001-00
	INCL	201001000
	IPCI	C10B0047-00 [ICM,7]; C10G0001-00 [ICS,7]
	NCL	201/001.000
	ECLA	C01B031/04; C10B055/00; H01M004/58E2
JP 2002083595	IPCI	H01M0004-58 [ICM,7]; C01B0031-00 [ICS,7]; C01B0031-04 [ICS,7]; C10B0057-04 [ICS,7]; C10C0003-02 [ICS,7]; H01M0004-02 [ICS,7]; H01M0010-40 [ICS,7]
EP 1186646	IPCI	C10C0003-00 [ICM,6]; C10B0055-00 [ICS,6];

C01B0031-04 [ICS,6]; H01M0004-58 [ICS,6];  
 H01M0010-40 [ICS,6]  
 ECLA C01B031/04; C10B055/00; H01M004/58E2  
 AB When a pitch compn., prepd. by mixing 100 wt. parts of  
 mesophase pitch with 10-1,000 wt. parts of  
 coal-tar pitch, is heat-treated at  $\geq 500^\circ$ , it  
 is possible to produce a high-d. coke at a high yield while avoiding  
 foaming of the mesophase pitch. In addn., when  
 the coke is graphitized at  $\geq 2,000^\circ$ , it is possible to  
 obtain an artificial graphite having a high graphitization  
 degree. Further, when the coke is first pulverized and then  
 graphitized at  $\geq 2,000^\circ$ , it is possible to obtain a  
 high-crystallinity graphite powder which can be suitably  
 used as a C material for a neg. electrode of non-aq.  
 solvent type secondary battery having a high discharge capacity and  
 a high charge-discharge efficiency.  
 ST coke manuf pitch; graphitization pitch coke; battery anode graphite  
 manuf  
 IT Fluoropolymers, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (as binder in manuf. of graphite anodes)  
 IT Coal tar pitch  
 Mesophase pitch  
 (in pitch coke manuf)  
 IT Battery anodes  
 (manuf. from graphitized pitch coke)  
 IT Graphitization  
 (of pitch coke)  
 IT Coke  
 RL: CPS (Chemical process); PEP (Physical, engineering or chemical  
 process); PROC (Process)  
 (pitch; manuf. from mesophase pitch and coal-tar pitch)  
 IT 24937-79-9, Polyvinylidene fluoride  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (as binder in manuf. of graphite anodes)  
 IT 91-20-3, Naphthalene, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (in manuf. of mesophase pitch)  
 IT 7782-42-5P, Graphite, preparation  
 RL: IMF (Industrial manufacture); PEP (Physical, engineering or  
 chemical process); PYP (Physical process); PREP (Preparation); PROC  
 (Process)  
 (manuf. from pitch coke)

L35 ANSWER 10 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2004:924152 HCAPLUS  
 DN 142:159400  
 ED Entered STN: 03 Nov 2004  
 TI Organic electrolytes for secondary lithium batteries with  
 carbon-based anodes  
 IN Do, Chil Hun; Mun, Seong In  
 PA Il Dong Chemical Co., Ltd., S. Korea; Korea Electro Technology  
 Research Institute  
 SO Repub. Korean Kongkae Taeho Kongbo, No pp. given  
 CODEN: KRXXA7  
 DT Patent  
 LA Korean  
 IC ICM H01M004-36  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
 Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI KR 2002048814 A 20020624 KR 2000-78088 20001218

PRAI KR 2000-78088

20001218

CLASS

PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES

KR 2002048814 ICM H01M004-36  
IPCI H01M0004-36 [ICM,7]

AB This org. electrolytic soln. consists of EC and PC for a Li battery with a **graphite**-based C anode. The battery has enhanced output power and increased energy d. due to a decrease in the irreversible sp. capacity of the assistant C **electrode**. The soln. consists of 0.8-33 vol.% propylene carbonate, 43-29 vol.% ethylene carbonate, and 56-38 vol.% di-Et carbonate, di-Me carbonate, ethyl-Me carbonate and methyl-Pr carbonate; and Li salt at 0.7-1.3M. When **mesophase** carbon microbeads (MCMB) is used as anode material, the electrolytic soln. comprises 0.8-20 vol.% PC, 45-35 vol.% EC, 56-45 vol.% di-Et carbonate. When the **pitch** coated **graphite** (PCG) is used as the anode material, the electrolytic soln. comprises 0.8-15 vol.% PC, 43-37 vol.% EC, 56-48 vol.% di-Et carbonate. When the hard C is used as the anode material, the electrolytic soln. comprises 0.8-20 vol.% PC, 45-35 vol.% EC, 56-45 vol.% di-Et carbonate.

ST org electrolyte carbon graphite anode lithium battery

IT Secondary batteries  
(lithium; org. electrolytes for secondary lithium batteries with carbon-based anodes)

IT Battery anodes  
Battery electrolytes  
(org. electrolytes for secondary lithium batteries with carbon-based anodes)

IT 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate  
108-32-7, Propylene carbonate 616-38-6, Dimethyl carbonate  
623-53-0, Ethyl-methyl carbonate 7440-44-0, Carbon, uses  
7782-42-5, Graphite, uses 56525-42-9, Methyl-propyl carbonate  
RL: DEV (Device component use); USES (Uses)  
(org. electrolytes for secondary lithium batteries with carbon-based anodes)

L35 ANSWER 11 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:913949 HCAPLUS

DN 142:97410

ED Entered STN: 01 Nov 2004

TI Coating of carbon cathode material for lithium secondary batteries

IN Han, Sang Cheol; Han, Yeong Su; Kang, Yong Muk; Lee, Jae Yeong; Park, Seong Cheol

PA Korea Advanced Institute of Science and Technology, S. Korea

SO Repub. Korean Kongkae Taeho Kongbo, No pp. given  
CODEN: KRXXA7

DT Patent

LA Korean

IC ICM H01M004-02

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
KR 2002010843	A	20020206	KR 2000-44367	

200007  
31

PRAI KR 2000-44367

20000731

## CLASS

PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES

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KR 2002010843 ICM H01M004-02  
IPCI H01M0004-02 [ICM,7]

AB This method for coating a highly cryst. C cathode material confers excellent surface characteristics which provides a battery with high energy d. and good performance. The highly cryst. C is coated with a pyrolytic C by pyrolysis of a hydrocarbon gas by tumbling CVD. The cryst. C material is selected from natural graphite, Kish graphite, SFG series, highly oriented pyrolytic graphite, mesophase pitch based C fiber (MPCF), and MCMB series and the hydrocarbon gas is selected from LPG, methane, butane, and benzene.

ST carbon cathode pyrolytic carbon coating lithium battery

IT Petroleum products  
(gases, liquefied; in pyrolytic carbon coating of carbon-based cathode material for lithium secondary batteries)

IT Secondary batteries  
(lithium; pyrolytic carbon coating of carbon-based cathode material for lithium secondary batteries)

IT Carbon fibers, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(mesophase pitch-based; pyrolytic carbon coating of carbon-based cathode material for lithium secondary batteries)

IT Battery cathodes  
(pyrolytic carbon coating of carbon-based cathode material for lithium secondary batteries)

IT 71-43-2, Benzene, processes 74-82-8, Methane, processes  
106-97-8, Butane, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(in pyrolytic carbon coating of carbon-based cathode material for lithium secondary batteries)

IT 7440-44-0, Carbon, uses 7782-42-5, Graphite, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(pyrolytic carbon coating of carbon-based cathode material for lithium secondary batteries)

L35 ANSWER 12 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:773879 HCAPLUS

DN 137:297338

ED Entered STN: 11 Oct 2002

TI Nonaqueous electrolyte secondary battery

IN Ota, Hideo; Kishi, Takashi

PA Toshiba Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 10 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01M010-40

ICS H01M010-40; H01M004-02; H01M004-58

CC 52-1 (Electrochemical, Radiational, and Thermal Energy Technology)  
Section cross-reference(s): 76

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 2002298915	A2	20021011	JP 2001-98184	



200103  
30

PRAI JP 2001-98184

20010330

## CLASS

PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES

JP 2002298915 ICM H01M010-40  
 ICS H01M010-40; H01M004-02; H01M004-58  
 IPCI H01M0010-40 [ICM,7]; H01M0010-40 [ICS,7];  
 H01M0004-02 [ICS,7]; H01M0004-58 [ICS,7]

AB The battery is characterized by having good discharge capacity, charging cycle life time, and is capable of avoiding cracking, firing, and sudden temp. increase when large elec. current occurs during nail sticking short circuit test. The battery comprises a pos. **electrode** having active material contg. lithium oxide and  $\geq 1$  metals selected from Al, Co, Fe, Mn, Ga, Ru, and Nb, a neg. **electrode**, and a nonaq. electrolyte substrate contg. phosphate ester and vinylene carbonate (such as ethylene carbonate and methylethyl carbonate).

ST nonaq electrolyte secondary battery active material

IT Esters, uses

RL: DEV (Device component use); USES (Uses)  
 (Phosphate; nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

IT **Mesophase pitch**

Secondary batteries

(nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

IT Carbon black, uses

Carbon fibers, uses

Fluoropolymers, uses

RL: TEM (Technical or engineered material use); USES (Uses)

(nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

IT 78-40-0, Triethyl phosphate 96-49-1, Ethylene carbonate

108-32-7, Propylene carbonate 512-56-1, Trimethyl phosphate

616-38-6, Dimethyl carbonate 623-53-0, Methylethylcarbonate

872-36-6, Vinylene carbonate 10463-05-5, Dimethylethylphosphate

RL: TEM (Technical or engineered material use); USES (Uses)

(nonaq. electrolyte secondary battery prepn.; nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

IT 12057-24-8, Lithium oxide, uses

RL: DEV (Device component use); USES (Uses)

(nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

IT 7439-96-5, Manganese, uses 7440-03-1, Niobium, uses 7440-18-8,

Ruthenium, uses 7440-48-4, Cobalt, uses 7440-55-3, Gallium, uses

RL: MOA (Modifier or additive use); TEM (Technical or engineered

material use); USES (Uses)

(nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

IT 7429-90-5, Aluminum, uses 7440-02-0, Nickel, uses

7440-50-8, Copper, uses 7782-42-5,

Graphite, uses 9002-88-4, Polyethylene 24937-79-9,

Polyvinylidene fluoride

RL: TEM (Technical or engineered material use); USES (Uses)

(nonaq. electrolyte secondary battery using phosphate ester and vinylene carbonate)

L35 ANSWER 13 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:253368 HCAPLUS

DN 136:265837

ED Entered STN: 05 Apr 2002  
 TI Graphite material for secondary lithium battery anode and its  
 manufacture  
 IN Okamura, Kaichiro; Takai, Yasuyuki; Sudo, Yoshinori  
 PA Petoca Ltd., Japan  
 SO Jpn. Kokai Tokkyo Koho, 8 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM H01M004-58  
 ICS C01B031-04; H01M004-02; H01M010-40  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
 Technology)

FAN.CNT 1	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2002100359	A2	20020405	JP 2000-291916	200009 26

PRAI JP 2000-291916 20000926

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2002100359	ICM	H01M004-58
	ICS	C01B031-04; H01M004-02; H01M010-40
	IPCI	H01M0004-58 [ICM,7]; C01B0031-04 [ICS,7]; H01M0004-02 [ICS,7]; H01M0010-40 [ICS,7]

AB The graphite has d002  $\leq 0.336$  nm, Lc  $\geq 100$  nm,  
 La  $\geq 100$  nm, and I100/I101  $> 1.6$  [I100 and I101 are the  
 intensities of diffraction peaks for its (100) and (101) faces] on  
 its x ray diffraction pattern. The graphite material is  
 prepd. by directly heating milled mesophase pitch  
 derived carbon fibers, by applying a current through the fibers via  
 a pair of electrodes, in a fluidized bed furnace with a  
 controlled atm.

ST lithium battery anode graphite manuf carbon fiber; cryst  
 characteristic graphite secondary lithium battery anode

IT Battery anodes  
 (cryst. structure and manuf. of graphite from carbon fibers for  
 secondary lithium battery anodes)

IT Carbon fibers, processes  
 RL: CPS (Chemical process); PEP (Physical, engineering or chemical  
 process); PROC (Process)  
 (milled mesophase pitch derived carbon fibers for manuf. of  
 graphite for secondary lithium battery anodes)

IT 7782-42-5P, Graphite, uses  
 RL: DEV (Device component use); IMF (Industrial manufacture); PREP  
 (Preparation); USES (Uses)  
 (cryst. structure and manuf. of graphite from carbon fibers for  
 secondary lithium battery anodes)

L35 ANSWER 14 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:183906 HCAPLUS

DN 136:234637

ED Entered STN: 15 Mar 2002

TI Nonaqueous electrolyte secondary battery for portable electronic  
 appliances

IN Satoh, Asako; Fujiwara, Masashi; Koiwa, Kaoru; Sekino, Masahiro;  
 Shimura, Nao; Hasebe, Hiroyuki; Oguchi, Masayuki; Onumai, Masayuki

PA Kabushiki Kaisha Toshiba, Japan

SO Eur. Pat. Appl., 39 pp.

CODEN: EPXXDW  
 DT Patent  
 LA English  
 IC ICM H01M010-40  
 ICS H01M004-58  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1187245	A2	20020313	EP 2001-307571	20010906
	EP 1187245	A3	20040421		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	US 2002055047	A1	20020509	US 2001-945808	20010905
	US 6852451	B2	20050208		
	JP 2002329529	A2	20021115	JP 2001-269299	20010905
	CN 1343022	A	20020403	CN 2001-132683	20010906
PRAI	JP 2000-270529	A	20000906		
	JP 2001-54938	A	20010228		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 1187245	ICM	H01M010-40
	ICS	H01M004-58
	IPCI	H01M0010-40 [ICM,6]; H01M0004-58 [ICS,6]
	ECLA	H01M004/58E2; H01M010/40L2
US 2002055047	IPCI	H01M0010-40 [ICM,7]; H01M0002-02 [ICS,7]
	NCL	429/337.000
	ECLA	H01M004/58E2; H01M010/40L2
JP 2002329529	IPCI	H01M0010-40 [ICM,7]; H01M0010-40 [ICS,7]; H01M0002-02 [ICS,7]; H01M0004-02 [ICS,7]; H01M0004-58 [ICS,7]
CN 1343022	IPCI	H01M0010-36; H01M0010-40

AB Disclosed is a nonaq. electrolyte secondary battery, comprising a case having a wall thickness not larger than 0.3 mm, a pos. electrode provided in the case, a neg. electrode provided in the case and the neg. electrode contg. a carbonaceous material capable of absorbing-desorbing lithium ions, and a nonaq. electrolyte provided in the case and the nonaq. electrolyte contg. a nonaq. solvent including  $\gamma$ -butyrolactone and a solute dissolved in the nonaq. solvent, wherein after being discharged to 3-V with a current of 0.2-C at room temp., the voltage redn. caused by the self-discharge at 60° is not larger than 1.5 V in 3 wk.

ST battery nonaq electrolyte portable electronic appliance use

IT Carbon fibers, uses

RL: DEV (Device component use); USES (Uses)  
 (mesophase pitch-based; nonaq. electrolyte secondary battery for portable electronic appliances)

IT Electric apparatus  
 Secondary batteries  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

IT Carbonaceous materials (technological products)  
 RL: DEV (Device component use); USES (Uses)  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

IT Carbon black, uses  
 Fluoropolymers, uses  
 RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses)  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

IT Mesophase pitch  
 (precursor for carbonaceous material; nonaq. electrolyte secondary battery for portable electronic appliances)

IT 96-49-1, Ethylene carbonate 7429-90-5, Aluminum, uses  
 7440-50-8, Copper, uses 9002-88-4, Polyethylene  
 12190-79-3, Cobalt lithium oxide colio2 14283-07-9, Lithium tetrafluoroborate  
 RL: DEV (Device component use); USES (Uses)  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

IT 96-48-0,  $\gamma$ -Butyrolactone 7782-42-5, Graphite  
 , uses 24937-79-9, PvdF  
 RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses)  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

IT 7439-93-2, Lithium, uses  
 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

IT 872-50-4, n-Methylpyrrolidone, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (nonaq. electrolyte secondary battery for portable electronic appliances)

L35 ANSWER 15 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:811781 HCAPLUS

DN 137:327379

ED Entered STN: 25 Oct 2002

TI Continuous production of trilaminates by coextrusion for polymer lithium batteries

IN Naarmann, Herbert; Kruger, Franz Josef; Schaefer, Tim

PA Dilo Trading A.-G., Switz.

SO Ger. Offen., 10 pp.

CODEN: GWXXBX

DT Patent

LA German

IC ICM H01M010-38

ICS H01M010-40

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

Section cross-reference(s): 38

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	
PI	DE 10118639	A1	20021024	DE 2001-10118639	20010412
PRAI	DE 2001-10118639		20010412		
CLASS					

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
DE 10118639	ICM	H01M010-38
	ICS	H01M010-40
	IPCI	H01M0010-38 [ICM,7]; H01M0010-40 [ICS,7]
	ECLA	H01M010/38; H01M010/40L2
AB	<p>The invention concerns the prodn. of Trilaminates, consisting of an anode composite, polymer electrolytes and a cathode composite, which are provided on the cathode side and on the anode side with a metallic grid. The prodn. is carried out continuously, preferably via coextrusion. The systems thus obtained form the basis for rechargeable polymer lithium batteries. The procedure according to invention contains the prodn. of anode masses, cathode material as well as the polymer gel electrolyte, which are: (1) homogeneously developed, (2) agree in structural viscosity and rheol., and (3) defined in shape by extrusion; and can be continuously formed as bands with reproducible wts. and laminated. The anode mass consists of graphite, preferably synthetic, e.g., mesocarbon microbeads with spherical particles or graphite fibers as well as a polymer binder e.g. polyfluoroelastomeres, polyolefins, polybutadiene or styrene copolymers, as well as polymethacrylates with alc. residues C4-C20, and polyvinyl compds. such as polyvinylpyrrolidone, polyvinylimidazole, polyvinylpyridin etc. and their copolymers, e.g. with methacrylic acid ester with alc. residues C4-C20, and a conducting salt e.g., LiPF6 or Li oxalato borates, etc.</p>	
ST	lithium secondary battery polymer trilaminate coextrusion	
IT	<p>Battery anodes          Battery cathodes          Extrusion of plastics and rubbers          Laminated materials          (continuous prodn. of trilaminates by coextrusion for polymer lithium batteries)</p>	
IT	<p>Fluoro rubber          Isoprene-styrene rubber          Polyolefins          RL: DEV (Device component use); USES (Uses)          (continuous prodn. of trilaminates by coextrusion for polymer lithium batteries)</p>	
IT	<p>Zeolites (synthetic), uses          RL: MOA (Modifier or additive use); USES (Uses)          (continuous prodn. of trilaminates by coextrusion for polymer lithium batteries)</p>	
IT	<p>Carbon fibers, uses          RL: DEV (Device component use); USES (Uses)          (graphite; continuous prodn. of trilaminates by coextrusion for polymer lithium batteries)</p>	
IT	<p>Secondary batteries          (lithium; continuous prodn. of trilaminates by coextrusion for polymer lithium batteries)</p>	
IT	<p>Battery electrolytes          (polymer gel; continuous prodn. of trilaminates by coextrusion for polymer lithium batteries)</p>	
IT	<p>96-49-1, Ethylene carbonate 108-32-7, Propylene carbonate          7782-42-5, Graphite, uses 7791-03-9, Lithium perchlorate          9003-17-2, Polybutadiene 9003-39-8, Polyvinylpyrrolidone          9003-47-8, Polyvinylpyridine 9003-53-6, Polystyrene 12031-65-1,          Lithium nickel oxide linio2 12057-17-9, Lithium manganese oxide          limn2o4 12190-79-3, Cobalt lithium oxide colio2 14283-07-9,          Lithium tetrafluoroborate 21324-40-3, Lithium hexafluorophosphate          25087-26-7D, Polymethacrylic acid, alkyl esters, with C4-20 alcs.</p>	

25232-42-2, Polyvinylimidazole 33454-82-9, Lithium triflate  
 37296-91-6, Lithium molybdenum oxide 37349-20-5, Lithium tungsten  
 oxide 39302-37-9, Lithium titanium oxide 473540-08-8  
 RL: DEV (Device component use); USES (Uses)  
 (continuous prodn. of trilaminates by coextrusion for polymer  
 lithium batteries)  
 IT 7631-86-9, Silica, uses 9011-17-0, Kynar 2801  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (continuous prodn. of trilaminates by coextrusion for polymer  
 lithium batteries)  
 IT 25038-32-8  
 RL: DEV (Device component use); USES (Uses)  
 (isoprene-styrene rubber, continuous prodn. of trilaminates by  
 coextrusion for polymer lithium batteries)  
 IT 7440-44-0, Carbon, uses  
 RL: DEV (Device component use); USES (Uses)  
 (mesocarbon microbeads; continuous prodn. of trilaminates by  
 coextrusion for polymer lithium batteries)  
 RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 RE  
 (1) Anon; DE 10020031 A1 HCAPLUS

L35 ANSWER 16 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 2  
 AN 2002:719538 HCAPLUS  
 DN 138:58831  
 ED Entered STN: 23 Sep 2002  
 TI Morphology/Behavior Relationship in Reversible Electrochemical  
 Lithium Insertion into Graphitic Materials  
 AU Aurbach, Doron; Teller, Hanan; Levi, Elena  
 CS Department of Chemistry, Bar-Ilan University, Ramat-Gan, 52900,  
 Israel  
 SO Journal of the Electrochemical Society (2002), 149(10), A1255-A1266  
 CODEN: JESQAN; ISSN: 0013-4651  
 PB Electrochemical Society  
 DT Journal  
 LA English  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 66, 72  
 AB Different types of **graphite** particles, including  
**synthetic** flakes, natural **graphite** flakes,  
**spherical graphite** particles (mesocarbon  
 microbeads), and combinations of graphite flakes and disordered C  
 particles were studied as **electrode** materials in Li  
 insertion/deinsertion processes. The morphol. of the graphite  
 particles plays an important role in their stability. One of the  
 key factors is the structure of the facets perpendicular to the  
 basal planes, through which Li insertion takes place. The smoother  
 these facets are, and the less crevices they contain, the more  
 stable are the graphite particles in Li insertion reactions. This  
 is attributed to the fact that the surface reactions on the  
 cathodically polarized graphite **electrodes** in the nonaq.  
 Li salt solns. involve gas formation, which may have a detrimental  
 effect on the integrity of the graphite particles. Another  
 important factor is the existence of some degree of disorder and  
 distortion in the graphite structure. More distortion in the  
 ordered graphite structure may lead to higher stability. The  
 detrimental effect of propylene carbonate in ethylene  
 carbonate-based solns. may lead to an explanation of the factors  
 that det. stability or destruction of lithiated graphite  
**electrodes**. The correlation between the morphol. of the  
 graphite particles and their orientation in practical  
**electrodes** is also discussed. Chronopotentiometry, electron

- microscopy, and x-ray diffraction were used in this study.
- ST graphite electrode morphol failure lithium electrochem insertion; battery graphite anode morphol failure lithium electrochem insertion
- IT Intercalation  
(electrochem.; morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries)
- IT Battery anodes  
(morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries)
- IT Intercalation  
(retro, electrochem.; morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries)
- IT 7782-42-5, Graphite, uses  
RL: DEV (Device component use); PRP (Properties); USES (Uses)  
(BSP 20, NG 15 and UF 8, synthetic and natural flakes, anodes; morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries)
- IT 7440-44-0, Carbon, uses  
RL: DEV (Device component use); PRP (Properties); USES (Uses)  
(anodes; morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries)
- IT 96-49-1, Ethylene carbonate 108-32-7, Propylene carbonate 616-38-6, Dimethyl carbonate  
RL: CPS (Chemical process); DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
(electrolyte contg.; morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries with)
- IT 7791-03-9, Lithium perchlorate ( $\text{LiClO}_4$ ) 21324-40-3, Lithium hexafluorophosphate ( $\text{LiPF}_6$ )  
RL: DEV (Device component use); USES (Uses)  
(electrolyte; morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries with)
- IT 7439-93-2, Lithium, processes  
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)  
(morphol./behavior relationship in reversible electrochem. lithium intercalation into graphitic anode materials for batteries)

RE.CNT 35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

- (1) Aurbach, D; Electrochim Acta 1994, V39, P2559
- (2) Aurbach, D; Electrochim Acta 1999, V45, P67 HCAPLUS
- (3) Aurbach, D; J Electrochem Soc 1996, V143, P3809 HCAPLUS
- (4) Aurbach, D; J Electrochem Soc 1996, V143, PL273
- (5) Aurbach, D; J Electrochem Soc 2001, V148, PA525
- (6) Aurbach, D; J Phys Chem B 1997, V101, P2195 HCAPLUS
- (7) Aurbach, D; J Power Sources 1993, V43, P47
- (8) Aurbach, D; J Power Sources 1999, V81-82, P95 HCAPLUS
- (9) Aurbach, D; Nonaqueous Electrochemistry, Chap 4 1999
- (10) Dahn, C; J Power Sources 2001, V101, P96
- (11) Dahn, J; Lithium Batteries, New Materials, Developments and Perspectives, Chap 1 1994
- (12) Fong, R; J Electrochem Soc 1990, V137, P2009 HCAPLUS
- (13) Gong, J; Carbon 1999, V37, P1409 HCAPLUS

- (14) Joho, F; J Power Sources 2001, V97-98, P78 HCAPLUS
- (15) Joint Committee Of Powder Diffraction Standards; JCPDS ICDD, File no 41-1487
- (16) Joint Committee Of Powder Diffraction Standards; JCPDS ICDD, File no 75-2078
- (17) Lampe-Onnerud, C; J Power Sources 2001, V97-98, P133 HCAPLUS
- (18) Naji, A; Carbon 1998, V36, P1347 HCAPLUS
- (19) Noel, M; J Power Sources 1998, V72, P53 HCAPLUS
- (20) Ohta, A; J Power Sources 1995, V54, P6 HCAPLUS
- (21) Oliver, J; J Power Sources 2001, V97-98, P151
- (22) Peled, E; Handbook of Battery Materials, Part III, Chap 6 1999
- (23) Peled, E; Lithium Batteries, Chap 3 1983
- (24) Rosoken, J; J Electrochem Soc 1996, V143, P2417
- (25) Tran, T; J Appl Electrochem 1996, V26, P1161 HCAPLUS
- (26) Winter, M; Adv Mater 1998, V10, P725 HCAPLUS
- (27) Winter, M; Handbook of Battery Materials, Part III, Chap 5 1999
- (28) Winter, M; J Electrochem Soc 1998, V145, P428 HCAPLUS
- (29) Wyckoff, R; Crystal Structures, 2nd ed 1965, V1, P27
- (30) Xing, W; J Electrochem Soc 1996, V143, P3482 HCAPLUS
- (31) Xue, J; J Electrochem Soc 1995, V142, P3668 HCAPLUS
- (32) Yang, C; J Power Sources 1998, V72, P66 HCAPLUS
- (33) Zaghib, K; J Power Sources 2001, V97-98, P97 HCAPLUS
- (34) Zanem, D; J Power Sources 2001, V97-98, P146
- (35) Zheng, T; Synth Met 1995, V73, P1 HCAPLUS
- L35 ANSWER 17 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN
- AN 2002:279785 HCAPLUS
- DN 136:372201
- ED Entered STN: 15 Apr 2002
- TI Electrochemical properties of MCMBs as anode for lithium ion battery
- AU Li, Baohua; Guo, Ruixia; Li, Kaixi; Lu, Chunxiang; Ling, Licheng
- CS State Key Laboratory of Coal Conversion, Institute of Coal Chemistry, Chinese Academy of Sciences, Taiyuan, 030001, Peop. Rep. China
- SO Preprints of Symposia - American Chemical Society, Division of Fuel Chemistry (2002), 47(1), 187-188
- CODEN: PSADPZ; ISSN: 1521-4648
- PB American Chemical Society, Division of Fuel Chemistry
- DT Journal
- LA English
- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
- AB Mesophase microbeads (MCMBs) prepd. from petroleum pitch or coal tar is one of the candidate carbon materials for the electrodes used to further increase the energy d. of lithium secondary batteries. In this regard, the charge and discharge properties of MCMBs with different mean diam. and size distribution were examd. The MCMBs heat-treated at 973 K were found to have a higher reversible capacity than the theor. lithium storage capacity of a graphite anode, suggesting that there is a cavity mechanism for the charge-discharge reaction. The mean diam. and size distribution of MCMBs were found to have an influence on the charge-discharge properties of MCMBs.
- ST mesocarbon microbead anode lithium ion battery
- IT Battery anodes  
(electrochem. properties of mesocarbon microbeads as anode for lithium-ion batteries)
- IT 7440-44-0, Carbon, uses  
RL: CPS (Chemical process); DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
(electrochem. properties of mesocarbon microbeads as anode for



lithium-ion batteries)

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

- (1) Bondarenko, G; Carbon 1998, V36(7-8), P1107 HCAPLUS
- (2) Buiel, E; J Electrochem Soc 1998, V145(6), P1977 HCAPLUS
- (3) Ishikawa, M; Journal of Power Sources 1996, V62, P229
- (4) Liu, Y; Carbon 1996, V34(2), P193 HCAPLUS
- (5) Mabuchi, A; J Electrochem Soc 1995, V142(4), P1041 HCAPLUS
- (6) Naji, A; Carbon 1995, V33(9), P1347
- (7) Pled, E; J Electrochem Soc 1996, V143(1), PL4

L35 ANSWER 18 OF 53 INSPEC (C) 2005 IEE on STN

AN 2002:7407826 INSPEC DN A2002-22-8245-018

TI Morphology/behavior relationship in reversible electrochemical lithium insertion into graphitic materials.

AU Aurbach, D.; Teller, H.; Levi, E. (Dept. of Chem., Bar-Ilan Univ., Ramat-Gan, Israel)

SO Journal of the Electrochemical Society (Oct. 2002) vol.149, no.10, p.A1255-66. 35 refs.

Doc. No.: S0013-4651(02)00110-6

Published by: Electrochem. Soc

Price: CCCC 0013-4651/2002/149(10)/1255/12/\$7.00

CODEN: JESQAN ISSN: 0013-4651

SICI: 0013-4651(200210)149:10L.A1255:MBRR;1-P

DT Journal

TC Experimental

CY United States

LA English

AB Different types of **graphite** particles, including**synthetic flakes, natural graphite flakes,****graphite fibers, round-shaped graphite****particles (mesocarbon microbeads), and combinations of graphite****flakes and disordered carbon particles were studied as****electrode materials in lithium insertion/deinsertion****processes. We suggest that the morphology of the graphite particles****plays an important role in their stability. One of the key factors****is the structure of the facets, perpendicular to the basal planes,****through which Li insertion takes place. The smoother these facets****are, and which contain fewer crevices, the more stable are the****graphite particles in Li insertion reactions. This is attributed to****the fact that the surface reactions of the cathodically polarized****graphite electrodes in the nonaqueous Li salt solutions****involve gas formation, which may have a detrimental effect on the****integrity of the graphite particles. Another important factor is the****existence of some degree of disorder and distortion in the graphite****structure. More distortion in the ordered graphite structure may****lead to higher stability. A clue to the understanding of the factors****that determine stability or destruction of lithiated graphite****electrodes is the pronounced detrimental effect of the****presence of propylene carbonate in ethylene carbonate-based****solutions, as discussed herein. Another important factor dealt with****in this paper is the correlation between the morphology of the****graphite particles and their orientation in practical****electrodes. The major tools for this study included****chronopotentiometry, electron microscopy, and X-ray diffraction.**

CC A8245 Electrochemistry and electrophoresis; A6865 Low-dimensional

CT ELECTROCHEMICAL **ELECTRODES**; ELECTRON MICROSCOPY; GRAPHITE

CT INTERCALATION COMPOUNDS; LITHIUM; X-RAY DIFFRACTION

ST morphology; reversible electrochemical lithium insertion; graphitic

material; graphite particles; synthetic flakes; natural graphite

flakes; graphite fibers; mesocarbon microbeads; disordered carbon

particles; electrode material; stability; facet structure;  
 surface reaction; cathodic polarization; nonaqueous Li salt  
 solution; disorder; gas formation; distortion; graphite structure;  
 propylene carbonate; ethylene carbonate; particle orientation;  
 chronopotentiometry; electron microscopy; X-ray diffraction; C-Li  
 CHI CLi bin, Li bin, C bin  
 ET Li; C\*Li; C-Li; CLi; C cp; cp; Li cp

L35 ANSWER 19 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2001:655019 HCAPLUS  
 DN 135:229339  
 ED Entered STN: 07 Sep 2001  
 TI Nonaqueous electrolyte secondary batteries with uniform discharging  
 end voltage  
 IN Tatebayashi, Yoshinao; Igasaki, Yoshiyuki; Kanda, Motoshi  
 PA Toshiba Corp., Japan  
 SO Jpn. Kokai Tokkyo Koho, 9 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM H01M004-02  
 ICS H01M004-02; H01M004-58; H01M010-40  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
 Technology)

FAN.CNT 1					
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
JP 2001243943	A2	20010907	JP 2000-54589	20000229	
JP 3705728	B2	20051012			
PRAI JP 2000-54589		20000229			

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2001243943	ICM	H01M004-02
	ICS	H01M004-02; H01M004-58; H01M010-40
	IPCI	H01M0004-02 [ICM,7]; H01M0004-02 [ICS,7]; H01M0004-58 [ICS,7]; H01M0010-40 [ICS,7]

AB The batteries comprise (A) cathode active material mixts. contg. Li  
 Ni mixed oxides and spinel-type Li Mn oxides and (B) carbonaceous  
 anode active material mixts. contg. nongraphitic carbon materials  
 and graphite. The blending ratio of the nongraphitic carbon and  
 graphite is controlled to keep the discharge potential change rate  
 (at the anode operating potential) to  $\leq 20$  mV/(m-Ah/g). The  
 batteries are esp. suitable for their use under serial connection.  
 ST nonaq electrolyte secondary battery carbonaceous anode; nongraphitic  
 carbon graphite blend battery anode; lithium mixed oxide cathode  
 secondary battery; nickel lithium oxide cathode secondary battery;  
 manganese lithium oxide cathode secondary battery  
 IT Carbon fibers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (graphite, mesophase pitch-based,  
 anodes; nonaq. secondary lithium batteries with lithium mixed  
 oxide blended cathodes and nongraphitic carbon-  
 graphite blended anodes)  
 IT Secondary batteries  
 (lithium; nonaq. secondary lithium batteries with lithium mixed  
 oxide blended cathodes and nongraphitic carbon-graphite blended  
 anodes)  
 IT Battery anodes

(nonaq. secondary lithium batteries with lithium mixed oxide blended cathodes and nongraphitic carbon-graphite blended anodes)

IT 7782-42-5, Graphite, uses  
 RL: DEV (Device component use); USES (Uses)  
 (anode; nonaq. secondary lithium batteries with lithium mixed oxide blended cathodes and nongraphitic carbon-graphite blended anodes)

IT 172922-65-5, Lithium manganese oxide (Li1.06Mn1.94O4) 358751-60-7, Aluminum cobalt lithium nickel oxide (Al0.08Co0.21LiNi0.71O2)  
 RL: DEV (Device component use); USES (Uses)  
 (cathode; nonaq. secondary lithium batteries with lithium mixed oxide blended cathodes and nongraphitic carbon-graphite blended anodes)

IT 7440-44-0, Carbon, uses  
 RL: DEV (Device component use); USES (Uses)  
 (nongraphitic, anode; nonaq. secondary lithium batteries with lithium mixed oxide blended cathodes and nongraphitic carbon-graphite blended anodes)

L35 ANSWER 20 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2001:85650 HCAPLUS

DN 134:149864

ED Entered STN: 06 Feb 2001

TI Method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode

IN Hatano, Hitomi; Nagayama, Katsuhiko; Eguchi, Kunihiro; Fukuda, Tsuneyoshi; Yutani, Satoshi

PA Kawasaki Steel Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C10C003-02

ICS C01B031-00; C01B031-02; C01B031-04; C10C003-08; H01M004-04

CC 51-19 (Fossil Fuels, Derivatives, and Related Products)

Section cross-reference(s): 52, 57

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2001031976	A2	20010206	JP 1999-208371	19990723

PRAI JP 1999-208371 19990723

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2001031976	ICM	C10C003-02
	ICS	C01B031-00; C01B031-02; C01B031-04; C10C003-08; H01M004-04
	IPCI	C10C0003-02 [ICM,7]; C01B0031-00 [ICS,7]; C01B0031-02 [ICS,7]; C01B0031-04 [ICS,7]; C10C0003-08 [ICS,7]; H01M0004-04 [ICS,7]

AB The method is carried out by extn. and removal of light components (e.g., pyridine) from pitch contg. meso-phase phase (obtained from polymn. of condensed polycyclic arom. hydrocarbon in the presence of catalyst) using cyclic compd. and/or polycyclic arom. hydrocarbon. The treated meso-phase pitch is further under thermal treatment at 450° for 3 h to obtain carbonaceous material suitable for manufg. graphite as electrode material in Li secondary battery.

ST carbonaceous material graphite hydrocarbon

- mesophase pitch; battery electrode**  
**graphite pitch treatment**  
 IT Polycyclic compounds  
 RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
 (arom. hydrocarbons; method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT Battery electrodes  
 Graphitization  
**Mesophase pitch**  
 Polymerization catalysts  
 Secondary batteries  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT Carbonaceous materials (technological products)  
 RL: FMU (Formation, unclassified); NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); FORM (Formation, nonpreparative); PROC (Process); USES (Uses)  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT Cyclic compounds  
 RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT Aromatic hydrocarbons, uses  
 RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)  
 (polycyclic; method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT 7637-07-2, Boron fluoride (BF<sub>3</sub>), uses 7664-39-3, Hydrogen fluoride, uses  
 RL: CAT (Catalyst use); NUU (Other use, unclassified); USES (Uses)  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT 7782-42-5P, Graphite, uses  
 RL: DEV (Device component use); IMF (Industrial manufacture); NUU (Other use, unclassified); PREP (Preparation); USES (Uses)  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT 91-20-3, Naphthalene, processes  
 RL: PEP (Physical, engineering or chemical process); PROC (Process)  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)
- IT 110-86-1, Pyridine, processes  
 RL: PEP (Physical, engineering or chemical process); REM (Removal or disposal); PROC (Process)  
 (method for treatment of meso-phase pitch to produce carbonaceous material for graphite used in battery electrode)

L35 ANSWER 21 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2001:152411 HCAPLUS

DN 134:181116

ED Entered STN: 02 Mar 2001

TI Flat nonaqueous electrolyte secondary battery with improvements in heavy loading discharge characteristics

IN Suzuki, Masami; Hayami, Muneto; Udagawa, Kazuo; Iiduka, Kazuo; Ishihara, Naomi; Hirahara, Satoshi; Sakai, Hirotaka; Yoda, Kiyoto; Shikota, Masataka

PA Toshiba Battery Co., Ltd., Japan

SO Eur. Pat. Appl., 51 pp.

CODEN: EPXXDW

DT Patent

LA English

IC ICM H01M010-40

ICS H01M002-02; H01M010-04

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

Section cross-reference(s): 55

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1079454	A2	20010228	EP 2000-117368	20000823
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2001068160	A2	20010316	JP 1999-240964	19990827
JP 2001068143	A2	20010316	JP 1999-241290	19990827
JP 2001143763	A2	20010525	JP 1999-327679	19991118
JP 2002008729	A2	20020111	JP 2000-183000	20000619
JP 2002008727	A2	20020111	JP 2000-183001	20000619
TW 504854	B	20021001	TW 2000-89116426	20000815
US 6521373	B1	20030218	US 2000-641267	20000817
HK 1035605	A1	20050916	HK 2001-106014	20010827
US 2003138693	A1	20030724	US 2002-318177	20021213
US 2005271938	A1	20051208	US 2005-176400	20050708
PRAI JP 1999-240964	A	19990827		
JP 1999-241290	A	19990827		
JP 1999-327679	A	19991118		
JP 2000-183000	A	20000619		
JP 2000-183001	A	20000619		
US 2000-641267	A3	20000817		
US 2002-318177	A3	20021213		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 1079454	ICM	H01M010-40
	ICS	H01M002-02; H01M010-04
	IPC1	H01M0010-40 [ICM,6]; H01M0002-02 [ICS,6]; H01M0010-04 [ICS,6]
	ECLA	H01M002/02B6B2; H01M002/02E2C; H01M002/12D2;

H01M010/04C2; H01M010/40L2

JP 2001068160 IPCI H01M0010-40 [ICM,7]  
 JP 2001068143 IPCI H01M0010-04 [ICM,7]; H01M0004-02 [ICS,7];  
 H01M0004-58 [ICS,7]; H01M0010-40 [ICS,7]  
 JP 2001143763 IPCI H01M0010-40 [ICM,7]; H01M0002-04 [ICS,7];  
 H01M0002-08 [ICS,7]; H01M0004-02 [ICS,7]  
 JP 2002008729 IPCI H01M0010-40 [ICM,7]; H01M0004-64 [ICS,7];  
 H01M0004-74 [ICS,7]  
 JP 2002008727 IPCI H01M0010-40 [ICM,7]  
 TW 504854 IPCI H01M0006-14 [ICM,7]; H01M0010-38 [ICS,7];  
 H01M0002-00 [ICS,7]  
 US 6521373 IPCI H01M0006-46 [ICM,7]  
 NCL 429/162.000; 429/094.000; 429/124.000;  
 429/127.000  
 ECLA H01M002/02B6B2; H01M002/02E2C; H01M002/12D2;  
 H01M010/04C2; H01M010/40L2

HK 1035605 IPCI H01M [ICM,7]  
 US 2003138693 IPCI H01M0002-02 [ICM,7]; H01M0002-08 [ICS,7];  
 H01M0004-58 [ICS,7]; H01M0010-40 [ICS,7];  
 H01M0004-66 [ICS,7]  
 NCL 429/162.000  
 ECLA H01M002/02B6B2; H01M002/02E2C; H01M002/12D2;  
 H01M010/04C2; H01M010/40L2

US 2005271938 IPCI H01M0002-08 [ICM,7]; H01M0010-04 [ICS,7];  
 H01M0004-58 [ICS,7]; H01M0010-40 [ICS,7];  
 H01M0002-04 [ICS,7]; H01M0002-02 [ICS,7]  
 NCL 429/185.000; 429/128.000; 429/162.000;  
 429/231.800; 429/329.000; 429/330.000;  
 429/176.000; 429/175.000

AB In a flat nonaq. electrolyte secondary cell comprising an electricity-generating element including at least a **cathode**, a separator and an anode and a nonaq. electrolyte in the inside of a **cathode** case, a plurality of **electrode** units each consisting of the **cathode** and the anode opposite to each another via the separator are laminated to form an **electrode** group, or an **electrode** unit in a sheet form consisting of the **cathode** and the anode opposite to each another via the separator is wound to form an **electrode** group, or a sheet-shape **cathode** is wrapped with the separator except for a part contacting at inner face of **cathode** case and a sheet-shaped anode is set on the sheet-shaped **cathode** in a right angled position each other and then these **cathode** and anode are bent alternately to form an **electrode** group, and the total sum of the areas of the opposing **cathode** and anode in this **electrode** group is larger than the area of the opening of an insulating gasket in a sealed portion in the **cathode** case or than the area of an opening in a sealed plate in a sealed portion in the **cathode** case, whereby the discharge capacity upon heavy-loading discharge is significantly increased as compared with the conventional cells. Accordingly, while the size of the cell is small, the discharge capacity is increased as described above, and thus it is possible to provide a highly utilizable flat nonaq. electrolyte secondary cell. Further, in the flat nonaq. electrolyte secondary cell, problems which may be caused by the increased discharge capacity in the cell can be solved by improving the solvent and supporting electrolyte for the electrolyte or by various improvements in the **cathode** and anode cases.

ST nonaq electrolyte secondary battery flat  
 IT Battery electrolytes  
 Secondary batteries  
 (flat nonaq. electrolyte secondary battery with improvements in

heavy loading discharge characteristics)

IT Carbonaceous materials (technological products)  
 RL: DEV (Device component use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT Carbon black, uses  
 RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT Fluoropolymers, uses  
 Styrene-butadiene rubber, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT Carbon fibers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (mesophase pitch, graphitized; flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT Fluoropolymers, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (tape; flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 96-48-0,  $\gamma$ -Butyrolactone 96-49-1, Ethylene carbonate  
 9002-88-4, Polyethylene 12190-79-3, Cobalt lithium oxide colio2  
 12597-68-1, Stainless steel, uses 14283-07-9 21324-40-3, Lithium  
 hexafluorophosphate 326594-36-9 326594-37-0 326594-38-1  
 326594-39-2 326594-40-5 326594-41-6  
 RL: DEV (Device component use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 7782-42-5, Graphite, uses  
 RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 7440-02-0, Nickel, uses 7440-50-8, Copper, uses  
 RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 872-50-4, n-Methylpyrrolidone, uses 9000-11-7, Cmc 24937-79-9, PvdF  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 7429-90-5, Aluminum, uses 7440-03-1, Niobium, uses 7440-32-6, Titanium, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (stainless steel contg.; flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 9003-55-8  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (styrene-butadiene rubber, flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

IT 9002-84-0, PtfE  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (tape; flat nonaq. electrolyte secondary battery with improvements in heavy loading discharge characteristics)

L35 ANSWER 22 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2001:627403 HCAPLUS  
 DN 135:307430  
 ED Entered STN: 30 Aug 2001  
 TI Mesophase coal tar pitch as an intermediate stage in production of  
 electrode coke  
 AU Robak, Zbigniew; Kubica, Krystyna  
 CS Inst. Chem. Przerobki Wegla, Zabrze, Pol.  
 SO Zeszyty Naukowe Politechniki Slaskiej, Chemia (2001), 142, 173-180  
 CODEN: ZNSCAM; ISSN: 0372-9494  
 PB Wydawnictwo Politechniki Slaskiej  
 DT Journal  
 LA Polish  
 CC 57-8 (Ceramics)  
 Section cross-reference(s): 51  
 AB Electrode cokes is routinely produced by delayed coking of  
 carefully chosen petroleum- or coal tar-derived feedstocks. The  
 process has its advantages and disadvantages, and to avoid some of  
 its difficulties other ways of transforming liq. **pitches**  
 into hard cokes are still under investigation. We tried to obtain a  
 highly anisotropic coke, with good graphitizability, by chamber  
 coking of a **mesophase pitch**. This material was  
 obtained from the model plant for continuous thermal treatment of  
 light tar **pitches**, and typically contained .apprx.50%  
**mesophase**. **Mesophase pitch** was  
 carbonized in the semi-tech. chamber coking plant at 550°C.  
 The obtained coke was investigated for its suitability for the  
**electrode** industry. The procedure simulates the typical  
 manufg. process for **graphite electrodes**, using  
 as a feedstock the "**mesophase coke**" and a std. petroleum  
 coke. It appeared that the "**mesophase coke**" gave  
 graphitized product showing properties comparable with that obtained  
 from com. Texas petroleum coke.  
 ST mesophase coal tar pitch feedstock electrode coke manuf  
 IT **Mesophase pitch**  
 (coal tar **pitch**, precursor; processing of  
**mesophase coal tar pitch** as feedstock for  
 prepn. of coke for **graphite electrode** manuf.)  
 IT Coke  
 RL: IMF (Industrial manufacture); PEP (Physical, engineering or  
 chemical process); PRP (Properties); PREP (Preparation); PROC  
 (Process)  
 (electrode; processing of **mesophase coal tar**  
**pitch** as feedstock for prepn. of coke for  
**graphite electrode** manuf.)  
 IT Coal tar **pitch**  
 (mesophase, precursor; processing of **mesophase**  
 coal tar **pitch** as feedstock for prepn. of coke for  
**graphite electrode** manuf.)  
 IT 7782-42-5, Graphite, miscellaneous  
 RL: MSC (Miscellaneous)  
 (electrodes; processing of **mesophase coal tar**  
**pitch** as feedstock for prepn. of coke for  
**graphite electrode** manuf.)

L35 ANSWER 23 OF 53 INSPEC (C) 2005 IEE on STN  
 AN 2003:7508108 INSPEC DN A2003-04-8630F-034; B2003-02-8410E-053  
 TI Development of new anodes for rechargeable lithium batteries.  
 AU Sandi, G. (Div. of Chem., Argonne Nat. Lab., IL, USA)  
 SO New Materials for Electrochemical Systems IV.. Extended Abstracts of  
 the Fourth International Symposium on New Materials for



## Electrochemical Systems

Editor(s): Savadogo, O.

Montreal, Que., Canada: Ecole Polytechnique de Montreal, 2001. p.162  
of xxiii+488 pp.

Conference: Montreal, Que., Canada, 9-13 July 2001

DT Conference Article

TC Practical

CY Canada

LA English

AB Summary form only given. Lithium ion secondary batteries are currently the best portable energy storage device for the consumer electronics market. The development of the lithium ion secondary batteries has been achieved by the use of selected carbon and **graphite** materials as an anode. The performance of lithium ion secondary batteries, such as the charge/discharge capacity, voltage profile and cyclic stability, depend strongly on the microstructure of the anode materials made of carbon and **graphite**. Due to the contribution of the carbon materials used in the anode in the last five years, the capacity of the typical Li ion battery has been improved 1.7 times. However, there are still active investigations to identify the key parameters of carbons that provide the improved anode properties, as carbon and **graphite** materials have large varieties in the microstructure, texture, crystallinity and morphology, depending on their preparation processes and precursor materials, as well as various forms such as powder, fibers and spherule. There is a strong correlation between the microstructural parameters and electrochemical properties of conventional and novel types of carbon materials for Li ion batteries, namely, graphitizable carbons such as milled **mesophase pitch**-based carbon fibers, polyparaphenylene based carbon heat-treated at low temperatures, boron doped graphitized materials, and templated carbons. a variety of tin oxide based compounds; SnO, SnO<sub>3</sub>, LiSnO<sub>3</sub>, and SnSiO<sub>3</sub> glass, have been tested as anode materials. These materials demonstrate discharge capacities on the order of 1000 mAh/(g Sn), which is consistent with the alloying capacity limit of 4.4 Li atoms per Sn atom, or 991 mAh/(g Sn). However, the irreversible capacities ranged from 200 mAh/(g active) to 700 mAh/(g active). By introducing lithium, lithium oxide and tin form first, which is then followed by the formation of the various LiSn alloy phases. Other investigations have focused on the **copper**-tin system at around the composition Cu<sub>6</sub>Sn<sub>5</sub> and have determined the effect on cycling and capacity of electrodes with various ratios of **copper** to tin. Other systems are compared and discussed in detail. The basic scientific theory can contribute to further developments of the Li ion batteries such as polymer batteries for consumer electronics, multimedia technology and future hybrid and electric vehicles.

CC A8630F Secondary cells; A8245 Electrochemistry and electrophoresis;  
B8410E Secondary cells

CT ANODES; CARBON; CARBON FIBRES; ELECTROCHEMICAL **ELECTRODES**;  
**GRAPHITE**; LITHIUM; SECONDARY CELLS

ST anodes; rechargeable lithium batteries; lithium ion secondary  
batteries; portable energy storage device; consumer electronics;  
**graphite materials**; carbon; anode; charge/discharge  
capacity; voltage profile; cyclic stability; microstructure; Li ion  
battery; texture; crystallinity; morphology; preparation processes;  
precursor materials; powder; fibers; spherule; microstructural  
parameters; electrochemical properties; Li ion batteries;  
graphitizable carbons; milled **mesophase pitch**-based carbon  
**fibers**; polyparaphenylene based carbon; boron doped graphitized  
materials; tin oxide based compounds; SnO; SnO<sub>3</sub>; LiSnO<sub>3</sub>; SnSiO<sub>3</sub>

glass; irreversible capacities; copper-tin system; Cu<sub>6</sub>Sn<sub>5</sub>;  
templated carbons; Li  
CHI Li int, Li el  
ET Li; O\*Sn; SnO; Sn cp; cp; O cp; SnO<sub>3</sub>; Li\*O\*Sn; Li sy 3; sy 3; O sy  
3; Sn sy 3; LiSnO<sub>3</sub>; Li cp; O\*Si\*Sn; Si sy 3; SnSiO<sub>3</sub>; Si cp; Sn;  
Li\*Sn; Li sy 2; sy 2; Sn sy 2; LiSn; Cu\*Sn; Cu sy 2; Cu<sub>6</sub>Sn<sub>5</sub>; Cu cp  
L35 ANSWER 24 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
AN 2002:143858 HCAPLUS  
DN 136:153932  
ED Entered STN: 25 Feb 2002  
TI Method for manufacturing cathode active material for lithium  
secondary battery  
IN Kim, Gwang Sik; Noh, Yeong Bae; Ryu, Jae Yul; Park, Jeong Jun; Woo,  
Il Gi; Yoon, Sang Yeong  
PA Samsung Sdi Co., Ltd., S. Korea  
SO Repub. Korean Kongkae Taeho Kongbo, No pp. given  
CODEN: KRXXA7  
DT Patent  
LA Korean  
IC ICM H01M004-48  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
Technology)

FAN.CNT 1  
PATENT NO. KIND DATE APPLICATION NO. DATE  
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PI KR 2000037532 A 20000705 KR 1998-52140 199812  
01

PRAI KR 1998-52140 19981201

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
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KR 2000037532	ICM	H01M004-48
	IPCI	H01M0004-48 [ICM,7]

AB A cathode active material for a lithium secondary battery  
is provided to enhance the charging and discharging characteristics  
and cycle life by manufg. the cathode active material by  
mixing a natural black lead with a synthetic  
black lead. A cathode active material  
for a lithium secondary battery is manufd. by mixing a natural  
black lead with a synthetic black  
lead. By mixing the natural black lead  
with the synthetic black lead, the difference of  
the elec. cond. between a-axis and c-axis is reduced by providing a  
conductive passage. The synthetic black lead is  
a milled carbon fiber made of a mesophase pitch.  
ST cathode active material lithium secondary battery  
IT Secondary batteries  
(lithium; method for manufg. cathode active material for lithium  
secondary battery)  
IT Battery cathodes  
Mesophase pitch  
(method for manufg. cathode active material for lithium secondary  
battery)  
IT Carbon fibers, uses  
RL: TEM (Technical or engineered material use); USES (Uses)  
(method for manufg. cathode active material for lithium secondary  
battery)  
IT 7439-92-1, Lead, uses  
RL: DEV (Device component use); USES (Uses)

(black, natural and synthetic; method for manufg. cathode active material for lithium secondary battery)

L35 ANSWER 25 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2000:53110 HCAPLUS

DN 132:110583

ED Entered STN: 23 Jan 2000

TI Secondary lithium batteries using cathodes containing mixed oxides

IN Ikasaki, Yoshiyuki; Sakurai, Katsuyuki; Hasebe, Hiroyuki

PA Toshiba Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 18 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01M004-62

ICS H01M004-02; H01M004-58; H01M010-40

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 2000021407	A2	20000121	JP 1998-184443	19980630

JP 2971451 B2 19991108  
 PRAI JP 1998-184443 19980630

#### CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2000021407	ICM	H01M004-62
	ICS	H01M004-02; H01M004-58; H01M010-40
	IPCI	H01M0004-62 [ICM,6]; H01M0004-02 [ICS,6]; H01M0004-58 [ICS,6]; H01M0010-40 [ICS,6]
AB		The batteries have cathodes contg. $\text{LiNi}_{1-x}\text{M}_x\text{O}_2$ ( $\text{M} = \text{element}; 0 < x \leq 0.5$ ), elec. conductors contg. carbonaceous materials (A) and (B) having av. sizes $\leq 100 \text{ nm}$ and $\geq 1 \mu\text{m}$ , resp., and binders contg. acrylic rubber copolymers and $\leq 45 \text{ wt.}\%$ (based on binders) vinylidene fluoride-type fluoropolymers, anodes, and nonaq. electrolyte solns. The batteries have long cycle life.
ST		battery cathode lithium oxide acrylic binder; carbon fluoropolymer lithium oxide battery cathode
IT		Synthetic rubber, uses RL: DEV (Device component use); USES (Uses) (acrylic acid-butadiene-itaconic acid-Me methacrylate-styrene; secondary Li batteries using cathodes contg. mixed oxides, C, and polymer binders for long cycle life)
IT		Synthetic rubber, uses RL: DEV (Device component use); USES (Uses) (acrylonitrile-butadiene-fumaric acid-itaconic acid-Me methacrylate-styrene; secondary Li batteries using cathodes contg. mixed oxides, C, and polymer binders for long cycle life)
IT		Synthetic rubber, uses RL: DEV (Device component use); USES (Uses) (acrylonitrile-butadiene-itaconic acid-Me methacrylate-styrene; secondary Li batteries using cathodes contg. mixed oxides, C, and polymer binders for long cycle life)
IT		Carbon fibers, uses RL: DEV (Device component use); USES (Uses) (graphite, mesophase pitch-based, anodes; secondary Li batteries using cathodes contg. mixed oxides, C, and polymer binders for long cycle life)

IT Secondary batteries  
 (lithium; secondary Li batteries using cathodes contg. mixed  
 oxides, C, and polymer binders for long cycle life)

IT Battery anodes  
 Battery cathodes  
 Battery electrolytes  
 (secondary Li batteries using cathodes contg. mixed oxides, C,  
 and polymer binders for long cycle life)

IT Carbon black, uses  
 Carbonaceous materials (technological products)  
 Fluoropolymers, uses  
 Oxides (inorganic), uses  
 RL: DEV (Device component use); USES (Uses)  
 (secondary Li batteries using cathodes contg. mixed oxides, C,  
 and polymer binders for long cycle life)

IT 75-05-8, Acetonitrile, uses 96-48-0,  $\gamma$ -Butyrolactone  
 96-49-1, Ethylene carbonate 105-58-8, Diethyl carbonate  
 108-32-7, Propylene carbonate 616-38-6, Dimethyl carbonate  
 623-53-0, Methyl ethyl carbonate 73506-93-1, Diethoxyethane  
 RL: DEV (Device component use); USES (Uses)  
 (electrolyte soln.; secondary Li batteries using cathodes contg.  
 mixed oxides, C, and polymer binders for long cycle life)

IT 21324-40-3, Lithium hexafluorophosphate  
 RL: DEV (Device component use); USES (Uses)  
 (electrolyte; secondary Li batteries using cathodes contg. mixed  
 oxides, C, and polymer binders for long cycle life)

IT 70857-13-5, Acrylonitrile-butadiene-itaconic acid-methyl  
 methacrylate-styrene copolymer 70857-14-6, Acrylic  
 acid-butadiene-itaconic acid-methyl methacrylate-styrene copolymer  
 118295-24-2, Acrylonitrile-butadiene-fumaric acid-itaconic  
 acid-methyl methacrylate-styrene copolymer  
 RL: DEV (Device component use); USES (Uses)  
 (rubber; secondary Li batteries using cathodes contg. mixed  
 oxides, C, and polymer binders for long cycle life)

IT 75-38-7D, Vinylidene fluoride, carboxy derivs., polymers  
 7782-42-5, Graphite, uses 24937-79-9, Poly(vinylidene fluoride)  
 RL: DEV (Device component use); USES (Uses)  
 (secondary Li batteries using cathodes contg. mixed oxides, C,  
 and polymer binders for long cycle life)

IT 113066-89-0P, Cobalt lithium nickel oxide (Co<sub>0.2</sub>LiNi<sub>0.8</sub>O<sub>2</sub>)  
 255063-53-7P, Aluminum cobalt lithium nickel oxide  
 (Al<sub>0.03</sub>Co<sub>0.17</sub>LiNi<sub>0.8</sub>O<sub>2</sub>) 255063-55-9P, Cobalt lithium nickel borate  
 oxide (Co<sub>0.17</sub>LiNi<sub>0.8</sub>(BO<sub>3</sub>)<sub>0.03</sub>O<sub>1.91</sub>)  
 RL: DEV (Device component use); PNU (Preparation, unclassified);  
 PREP (Preparation); USES (Uses)  
 (secondary Li batteries using cathodes contg. mixed oxides, C,  
 and polymer binders for long cycle life)

L35 ANSWER 26 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 2000:723263 HCAPLUS  
 DN 133:269460  
 ED Entered STN: 13 Oct 2000  
 TI Lithium secondary battery  
 IN Sato, Asako; Kuboki, Takashi; Yamada, Shuji; Hasebe, Hiroyuki;  
 Takami, Norio; Ohsaki, Takahisa; Kanda, Motoya  
 PA Kabushiki Kaisha Toshiba, Japan  
 SO Eur. Pat. Appl., 52 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA English  
 IC ICM H01M002-02  
 ICS H01M010-04

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1043782	A2	20001011	EP 2000-106079	20000330
EP 1043782	A3	20011114		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
US 6589690	B1	20030708	US 2000-536685	20000328
CN 1273439	A	20001115	CN 2000-118175	20000330
CN 1133228	B	20031231		
JP 2000348776	A2	20001215	JP 2000-94680	20000330
JP 3721044	B2	20051130		
TW 451511	B	20010821	TW 2000-89105965	20000330
JP 2004047479	A2	20040212	JP 2003-285821	20030804
PRAI JP 1999-89173	A	19990330		
JP 1999-89175	A	19990330		
JP 2000-94680	A3	20000330		

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 1043782	ICM	H01M002-02
	ICS	H01M010-04
	IPCI	H01M0002-02 [ICM,6]; H01M0010-04 [ICS,6]
	ECLA	H01M002/02E2D; H01M010/04; H01M010/04F
US 6589690	IPCI	H01H0069-02 [ICM,7]; H01M0006-00 [ICS,7]; H01M0004-58 [ICS,7]
	NCL	429/162.000; 029/623.100; 029/623.300; 029/623.500; 429/160.000; 429/164.000; 429/231.800; 429/231.950
	ECLA	H01M002/02E2D; H01M010/04; H01M010/04F
CN 1273439	IPCI	H01M0010-40 [ICM,7]; H01M0010-36 [ICS,7]
JP 2000348776	IPCI	H01M0010-40 [ICM,7]; H01M0010-40 [ICS,7]; H01M0002-02 [ICS,7]; H01M0002-18 [ICS,7]; H01M0010-04 [ICS,7]; H01M0004-58 [ICS,7]
TW 451511	IPCI	H01M0010-40 [ICM,7]
JP 2004047479	IPCI	H01M0010-40 [ICM,7]; H01M0002-02 [ICS,7]; H01M0004-02 [ICS,7]; H01M0004-58 [ICS,7]
	FTERM	5H011/AA03; 5H011/AA13; 5H011/CC02; 5H011/CC06; 5H011/CC10; 5H011/DD01; 5H011/DD03; 5H011/DD13; 5H011/KK01; 5H029/AJ05; 5H029/AJ12; 5H029/AK02; 5H029/AK03; 5H029/AK05; 5H029/AL01; 5H029/AL02; 5H029/AL03; 5H029/AL04; 5H029/AL06; 5H029/AL07; 5H029/AL08; 5H029/AL12; 5H029/AM02; 5H029/AM03; 5H029/AM04; 5H029/AM05; 5H029/AM07; 5H029/AM16; 5H029/BJ04; 5H029/BJ14; 5H029/DJ02; 5H029/DJ04; 5H029/EJ04; 5H029/EJ12; 5H029/HJ04; 5H050/AA07; 5H050/AA15; 5H050/BA17; 5H050/CA02; 5H050/CA05; 5H050/CA07; 5H050/CA08; 5H050/CA09; 5H050/CA11;

5H050/CB01; 5H050/CB02; 5H050/CB03; 5H050/CB05;  
 5H050/CB07; 5H050/CB08; 5H050/CB09; 5H050/CB12;  
 5H050/DA04; 5H050/DA19; 5H050/EA09; 5H050/EA24;  
 5H050/FA05; 5H050/HA04

- AB A secondary battery comprises an electrode group with a pos. electrode, a neg. electrode having a neg. electrode collector and a neg. electrode layer held to the collector, and a separator interposed between the pos. electrode and the neg. electrode layer, a nonaq. electrolyte held by the electrode group, and a jacket housing the electrode group and having a thickness of not more than 0.3 mm. The pos. electrode, the neg. electrode and the separator are formed integral, and the peeling strength between the neg. electrode layer and the separator is lower than the peeling strength between the neg. electrode layer and the neg. electrode collector.
- ST lithium secondary battery
- IT Fluoropolymers, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (binder; lithium secondary battery)
- IT Battery anodes  
 Battery cathodes  
 (lithium secondary battery)
- IT Carbon fibers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (lithium secondary battery)
- IT Carbon black, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (lithium secondary battery)
- IT Secondary batteries  
 (lithium; lithium secondary battery)
- IT Carbon fibers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (mesophase pitch-based; lithium secondary battery)
- IT Carbon fibers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (pitch-based; lithium secondary battery)
- IT Polyimides, uses  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (resin; lithium secondary battery)
- IT 25014-41-9, Polyacrylonitrile  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (adhesive; lithium secondary battery)
- IT 24937-79-9, PvdF  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (binder; lithium secondary battery)
- IT 96-48-0,  $\gamma$ -Butyrolactone 96-49-1, Ethylene carbonate  
 105-58-8, Diethyl carbonate 108-32-7, Propylene carbonate  
 7429-90-5, Aluminum, uses 7440-50-8, Copper,  
 uses 9002-88-4, Polyethylene 14283-07-9, Lithium  
 tetrafluoroborate 21324-40-3, Lithium hexafluorophosphate  
 111706-40-2, Cobalt lithium oxide CoLiO-102 113066-89-0, Cobalt  
 lithium nickel oxide Co<sub>0.2</sub>LiNi<sub>0.8</sub>O<sub>2</sub>  
 RL: DEV (Device component use); USES (Uses)  
 (lithium secondary battery)
- IT 7782-42-5, Graphite, uses  
 RL: MOA (Modifier or additive use); USES (Uses)  
 (lithium secondary battery)
- IT 9003-07-0, Polypropylene  
 RL: TEM (Technical or engineered material use); USES (Uses)  
 (lithium secondary battery)

IT 7440-44-0, Carbon, uses  
RL: DEV (Device component use); USES (Uses)  
(spheres or granules; lithium secondary battery)

L35 ANSWER 27 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
AN 1999:465067 HCAPLUS  
DN 131:288727  
ED Entered STN: 29 Jul 1999  
TI Characteristics of coke carbon modified with mesophase-pitch as a  
negative electrode for lithium ion batteries  
AU Sato, Yuichi; Kikuchi, Yasuo; Nakano, Takeshi; Okuno, Gaku;  
Kobayakawa, Koichi; Kawai, Takanobu; Yokoyama, Akira  
CS Faculty of Engineering, Department of Applied Chemistry, Kanagawa  
University, Kanagawa-ku, Yokohama, Japan  
SO Journal of Power Sources (1999), 81-82, 182-186  
CODEN: JPSODZ; ISSN: 0378-7753  
PB Elsevier Science S.A.  
DT Journal  
LA English  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
Technology)  
Section cross-reference(s): 57, 72  
AB To increase the charge-discharge capacity of carbon electrodes for  
lithium ion secondary batteries, coke carbon, a relatively cheap  
material, was modified with mesophase-pitch carbon by a heat  
treatment. While coke carbon powder, mesophase-pitch, and a mixt.  
thereof (4:1 by wt.) supplied between 0 and 1.5 V vs. Li/Li+ an  
initial discharge capacity of about 295 mA-h/g, 310 mA-h/g, and 310  
mA-h/g, resp., the modified coke deintercalated 400 mA-h/g of  
lithium with a high degree of reversibility. The difference in  
capacity between the modified carbon and mixt. are discussed based  
on the shape of their current-potential curves and their  
galvanostatic charge-discharge curves.  
ST lithium battery anode coke carbon  
IT Battery anodes  
Cyclic voltammetry  
(characteristics of coke carbon modified with mesophase-pitch as  
a neg. electrode for lithium ion batteries)  
IT Carbonaceous materials (technological products)  
Coke  
RL: DEV (Device component use); USES (Uses)  
(characteristics of coke carbon modified with mesophase-pitch as  
a neg. electrode for lithium ion batteries)  
IT Secondary batteries  
(lithium; characteristics of coke carbon modified with  
mesophase-pitch as a neg. electrode for lithium ion batteries)  
IT 7440-44-0, Carbon, uses 7782-42-5, Graphite,  
uses  
RL: DEV (Device component use); USES (Uses)  
(characteristics of coke carbon modified with mesophase  
-pitch as a neg. electrode for lithium ion  
batteries)

RE.CNT 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD  
RE  
(1) Alamgir, M; J Electrochem Soc 1993, V141, PL143  
(2) Bar-Tow, D; The Electrochemical Soc Meeting Abstracts 1996, V96-2,  
P1028  
(3) Besenhard, J; J Power Sources 1995, V54, P228 HCAPLUS  
(4) Disma, E; J Electrochem Soc 1996, V143, P3959  
(5) Eli, Y; J Electrochem Soc 1997, V144, P2968  
(6) Fong, R; J Electrochem Soc 1990, V137, P2009 HCAPLUS  
(7) Hara, M; J Phys Chem 1995, V99, P16338 HCAPLUS

- (8) Inagaki, M; Tanso (Carbon) 1984, V118, P165 HCAPLUS
- (9) JSPS; Tanso 1963, V36, P25
- (10) Kikuchi, M; J Electroanal Chem 1995, V396, P451 HCAPLUS
- (11) Kinoshita, K; Carbon Electrochemical and Physicochemical Properties 1988
- (12) Mabuchi, A; J Electrochem Soc 1995, V142, P1041 HCAPLUS
- (13) Nakajima, T; Denki Kagaku 1996, V64, P922 HCAPLUS
- (14) Ohsaki, T; Proceeding of the 34th Battery Symposium in Japan 1993, P79
- (15) Omaru, A; The Electrochemical Soc Extended Abstracts 1992, V92-2, P34
- (16) Sato, K; Science 1994, V264, P556 HCAPLUS
- (17) Sonbe, N; Proceedings of the 35th Battery Symposium in Japan 1994, P47
- (18) Takami, N; J Electrochem Soc 1995, V142, P371 HCAPLUS
- (19) Tatsumi, K; J Electrochem Soc 1995, V142, P716 HCAPLUS
- (20) Tuinstra, F; J Chem Phys 1970, V53, P1126 HCAPLUS
- (21) Yata, S; Synth Met 1994, V62, P153 HCAPLUS

L35 ANSWER 28 OF 53 JAPIO (C) 2005 JPO on STN  
 AN 1998-121054 JAPIO  
 TI PRODUCTION OF NONSCALELIKE CARBONACEOUS POWDER AND GRAPHITE POWDER  
 IN SAKAI YUKIO; FUJIURA TAKATSUGU  
 PA MITSUBISHI GAS CHEM CO INC  
 PI JP 10121054 A 19980512 Heisei  
 AI JP 1996-278012 (JP08278012 Heisei) 19961021  
 PRAI JP 1996-278012 19961021  
 SO PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined Applications, Vol. 1998

IC ICM C10C003-02  
 ICS C01B031-02; C01B031-04; C10C003-04; C10C003-10; H01M004-02; H01M004-04; H01M004-58

AB PROBLEM TO BE SOLVED: To obtain nearly **spherical** nonscalelike carbonaceous powder by heat-treating specified **synthetic mesophase** pitch in a specified temperature range in nonoxidizing atmosphere and grinding the treated pitch.  
 SOLUTION: This powder is obtained by heat-treating **mesophase** pitch obtained by polymerizing fused polycyclic hydrocarbon or a material containing it in a temperature region of 470-700°C in the presence of hydrogen fluoride or boron trifluoride in nonoxidizing atmosphere and grinding the treated pitch. The nonoxidizing gas used is exemplified by nitrogen or argon. The powder is graphitized at 1,900-3,000°C without undergoing fusion among particles to obtain nonscalelike higher crystalline **graphite** powder retaining the shape after the grinding. The **graphite** powder is desirable as a negative **electrode** material of a lithium ion battery and is very advantageous to improve **electrode** packing properties and to thereby attain a high energy density. Further, it has excellent conductivity and therefore it is usable also as a conductive filler for various materials.  
 COPYRIGHT: (C)1998,JPO

L35 ANSWER 29 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 1997:388601 HCAPLUS  
 DN 127:20944  
 ED Entered STN: 21 Jun 1997  
 TI Method for manufacture of carbonaceous material for nonaqueous lithium secondary batteries  
 IN Nishimura, Kasuke  
 PA Petoca, Ltd., Japan



SO Jpn. Kokai Tokkyo Koho, 9 pp.  
CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01M004-58

ICS D01F009-145; H01M004-02; H01M004-04; D21H013-50  
CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 09092283	A2	19970404	JP 1995-263710	19950919

PRAI JP 1995-263710

19950919

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 09092283	ICM	H01M004-58
	ICS	D01F009-145; H01M004-02; H01M004-04; D21H013-50
	IPCI	H01M0004-58 [ICM,6]; D01F0009-145 [ICS,6]; H01M0004-02 [ICS,6]; H01M0004-04 [ICS,6]; D21H0013-50 [ICS,6]
AB		The carbonaceous material is paper sheets 50-200 $\mu$ m thick having an apparent d. 1.0-1.5 g/cm <sup>3</sup> , prepd. from 5-30% C fibers 1-20 $\mu$ m in length and 5-10% mesophase pitch-based micro C beads and/or powd. graphite (natural or synthetic), and is used for cathodes of nonaq. Li secondary batteries.
ST		carbon fiber micro bead battery cathode
IT		Battery cathodes (carbon fibers and mesophase pitch-based beads for nonaq. lithium secondary batteries)
IT		Carbon fibers, uses RL: TEM (Technical or engineered material use); USES (Uses) (carbon fibers and mesophase pitch-based beads for nonaq. lithium secondary batteries)
IT		Secondary batteries (lithium; carbon fibers and mesophase pitch-based beads for nonaq. lithium secondary batteries)

L35 ANSWER 30 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1997:723810 HCAPLUS

DN 128:24881

ED Entered STN: 17 Nov 1997

TI High-capacity lithium-ion cells using graphitized mesophase-pitch-based carbon fiber anodes

AU Ohsaki, Takahisa; Kanda, Motoya; Aoki, Yoshiyasu; Shiroki, Hiroyuki; Suzuki, Shintaro

CS Saiwai-ku, 72 Horikawa-cho, Toshiba Corporation, Materials and Devices Research Laboratories, R+D Center, Kawasaki 210, Japan

SO Journal of Power Sources (1997), 68(1), 102-105  
CODEN: JPSODZ; ISSN: 0378-7753

PB Elsevier

DT Journal

LA English

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

AB The authors developed high-capacity lithium-ion cells using graphitized mesophase-pitch-based carbon fiber (MCF) as an anode material. The graphitized MCF is a highly graphitized carbon fiber with a radial-like texture in the cross

section. This structure contributes to the rapid diffusion of lithium ions inside the carbon fiber. The diffusion coeff. of lithium ions in the graphitized MCF was one order of magnitude larger than that for graphite, resulting in an excellent high-rate performance of the carbon electrode. The graphitized MCF anode showed larger capacity, higher rate capability, and better reversibility than the graphite anode. Prismatic cell (8.6 mm + 34 mm + 48 mm) with the graphitized MCF anode exhibited a large capacity of >1000 mA-h. At 3 A discharge, the prismatic cell had 95% of its capacity at 0.5 A discharge with a mid-discharge voltage of 3.35 V. The cell maintained >85% of its initial capacity after 500 cycles and showed high capacity at -20°. It has thus been demonstrated that the prismatic cell using the graphitized MCF anode has excellent performance, and is an attractive choice for the power sources of cellular phones and other appliances.

ST lithium ion battery development; carbon fiber graphitized anode  
lithium battery

IT Battery anodes  
Secondary batteries  
(development and performance of high-capacity lithium-ion  
batteries with graphitized mesophase-pitch-based carbon fiber  
anodes)

IT Carbon fibers, uses  
Carbon fibers, uses  
RL: DEV (Device component use); USES (Uses)  
(graphite, pitch-based; development and performance of  
high-capacity lithium-ion batteries with graphitized  
mesophase-pitch-based carbon fiber anodes)

RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD

- RE  
(1) Fong, R; J Electrochem Soc 1990, V137, P2009 HCAPLUS  
(2) Imanishi, N; J Electrochem Soc 1993, V140, P315 HCAPLUS  
(3) Inada, K; Proc Primary and Secondary Ambient Temperature Lithium  
Batteries 1988, V88-6, P530 HCAPLUS  
(4) Mohri, M; J Power Sources 1989, V26, P545 HCAPLUS  
(5) Sato, M; Primary and Secondary Lithium Batteries 1991, V91-3, P407  
HCAPLUS  
(6) Takami, N; J Electrochem Soc 1995, V142, P2564 HCAPLUS  
(7) Takami, N; J Electrochem Soc 1995, V142, P371 HCAPLUS  
(8) Yazami, R; J Power Sources 1983, V9, P365 HCAPLUS

L35 ANSWER 31 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 3  
AN 1995:503423 HCAPLUS

DN 122:244195

ED Entered STN: 22 Apr 1995

TI Nonaqueous electrolyte secondary lithium batteries

IN Sugimoto, Toyoji; Ozaki, Yoshuki; Ootsuka, Atsushi; Takai, Tooru

PA Matsushita Electric Ind Co Ltd, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01M010-40

ICS H01M004-02

CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
Technology)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 07037618	A2	19950207	JP 1993-181141	

PI JP 07037618

199307  
22

PRAI JP 1993-181141

19930722

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES		
JP 07037618	ICM	H01M010-40		
	ICS	H01M004-02		
	IPCI	H01M0010-40 [ICM,6]; H01M0004-02 [ICS,6]		
AB	The batteries comprise Li-contg. mixed oxide cathodes, nonaq. electrolytes, rechargeable cathodes of composite carbon materials consisting of spherical graphite particles having optically anisotropic single phase, and other graphite fine particles having an av. diam. smaller than that of another graphite particles. The spherical graphite particles may be manufd. by graphitization of mesocarbon microbeads from mesophase small spheres formed at carbonization of pitch, have lamella structures, and lattice distance (d002) 3.36-3.39 Å. The composite carbon materials may contain ≤20%, preferably 3-15%, the graphite fine particles to the spherical graphite particles. The graphite fine particles may be natural or synthetic graphite from petroleum or coal pitch.			
ST	graphite lithium battery anode			
IT	Anodes (battery, anodes contg. different graphite particles for secondary Li batteries for cycle performance)			
IT	Pitch (coal-tar, graphite from; anodes contg. different graphite particles for secondary Li batteries for cycle performance)			
IT	Pitch (petroleum, graphite from; anodes contg. different graphite particles for secondary Li batteries for cycle performance)			
IT	7782-42-5, Graphite, uses RL: DEV (Device component use); USES (Uses) (anodes contg. different graphite particles for secondary Li batteries for cycle performance)			
IT	7440-44-0, Mesocarbon microbeads, uses RL: DEV (Device component use); USES (Uses) (graphitized; anodes contg. different graphite particles for secondary Li batteries for cycle performance)			
L35	ANSWER 32 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN			
AN	1995:709126 HCAPLUS			
DN	123:88518			
ED	Entered STN: 29 Jul 1995			
TI	Secondary nonaqueous electrolyte batteries having improved anodes			
IN	Ootsuka, Atsushi; Sugimoto, Toyoji; Ozaki, Yoshuki; Takai, Tooru			
PA	Matsushita Electric Ind Co Ltd, Japan			
SO	Jpn. Kokai Tokyo Koho, 7 pp. CODEN: JKXXAF			
DT	Patent			
LA	Japanese			
IC	ICM H01M004-02 ICS H01M004-58; H01M010-40			
CC	52-2 (Electrochemical, Radiational, and Thermal Energy Technology)			
FAN.CNT 1				
	PATENT NO.	KIND DATE APPLICATION NO. DATE		
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PI JP 07134988 A2 19950523 JP 1993-281159 199311  
10

JP 3048808 B2 20000605  
PRAI JP 1993-281159 19931110

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 07134988	ICM	H01M004-02
	ICS	H01M004-58; H01M010-40
	IPCI	H01M0004-02 [ICM,6]; H01M0004-58 [ICS,6]; H01M0010-40 [ICS,6]

AB In the batteries having Li-contg. oxides as cathode active mass, powd. graphite-based rechargeable anodes, and nonaq. electrolytes, the graphite powders as Li-intercalatable and optically anisotropic spheres are obtained by graphitizing mesophase small spheres formed in low-temp. heating of pitches and have monophase lamellar structure, 002 plane lattice spacing (d002) by wide-angle x-ray diffraction 3.36-3.40 Å, and sp. surface area by BET method 0.7-5.0 m<sup>2</sup>/g. The batteries have high capacity, energy d., and storage property.

ST battery anode graphite powder

IT Anodes  
(battery, nonaq. electrolyte batteries using anodes of Li-intercalatable graphite powders having limited structure and surface)

IT 7439-93-2, Lithium, uses  
RL: DEV (Device component use); USES (Uses)  
(nonaq. electrolyte batteries using anodes of Li-intercalatable graphite powders having limited structure and surface)

IT 7782-42-5, Graphite, uses  
RL: DEV (Device component use); USES (Uses)  
(powd.; nonaq. electrolyte batteries using anodes of Li-intercalatable graphite powders having limited structure and surface)

L35 ANSWER 33 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1995:899204 HCAPLUS

DN 123:291865

ED Entered STN: 07 Nov 1995

TI Cathode for secondary lithium batteries and its manufacture

IN Takami, Norio; Ohsaki, Takahisa; Tamaki, Toshio; Nakajima, Hideyuki; Katsuta, Yasushi

PA Kabushiki Kaisha Toshiba, Japan; Petroca, Ltd.

SO Eur. Pat. Appl., 17 pp.  
CODEN: EPXXDW

DT Patent

LA English

IC ICM H01M004-58

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

## FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 675555	A1	19951004	EP 1995-104869	199503 31
EP 675555 R: DE, FR, GB	B1	19990728		
CN 1113351	A	19951213	CN 1995-103839	199503

31

CN 1048357 B 20000112  
JP 08069798 A2 19960312 JP 1995-97691

199503  
31

JP 3617550 B2 20050209  
US 5795678 A 19980818 US 1995-414195

199503  
31

PRAI JP 1994-85246 A 19940401

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 675555	ICM	H01M004-58
	IPCI	H01M0004-58 [ICM,6]
	ECLA	H01M004/58E2
CN 1113351	IPCI	H01M0004-62 [ICM,5]; H01M0004-64 [ICS,5]; H01M0004-38 [ICS,5]; H01M0004-04 [ICS,5]
JP 08069798	IPCI	H01M0004-58 [ICM,6]; D01F0009-145 [ICS,6]; H01M0004-02 [ICS,6]; H01M0004-04 [ICS,6]; H01M0010-40 [ICS,6]
US 5795678	IPCI	H01M0004-58 [ICM,6]; H01M0004-02 [ICS,6]
	NCL	429/218.100; 429/209.000
	ECLA	H01M004/58E2

AB The cathode comprises milled graphite fibers derived from mesophase pitch each having circumferential, upper end and lower end faces. Each milled graphite fiber is composed of graphite layers having voids between them as inlets and outlets for Li ions, and the circumferential, upper end and lower end faces having openings of the voids between the graphite layers. This cathode can be used to provide a secondary nonaq.-electrolyte battery of large charge and discharge capacities and high charge or discharge c.d.

ST graphite fiber milled battery cathode;  
pitch mesophase milled graphite fiber cathode

IT Cathodes  
(battery, milled graphite fibers from mesophase pitch for)

IT Carbon fibers, uses  
RL: DEV (Device component use); PNU (Preparation, unclassified); PRP (Properties); PREP (Preparation); USES (Uses)  
(graphite, cathodes for secondary lithium batteries)

IT Pitch  
(mesophase, battery cathodes from milled graphite fibers from)

L35 ANSWER 34 OF 53 JAPIO (C) 2005 JPO on STN

AN 1995-226204 JAPIO

TI MANUFACTURE OF NONAQUEOUS ELECTROLYTE SECONDARY BATTERY

IN OZAKI YOSHIYUKI; KOSHINA HIDE

PA MATSUSHITA ELECTRIC IND CO LTD

PI JP 07226204 A 19950822 Heisei

AI JP 1994-16338 (JP06016338 Heisei) 19940210

PRAI JP 1994-16338 19940210

SO PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined Applications, Vol. 1995

IC ICM H01M004-58  
ICS H01M004-02; H01M004-04; H01M004-96; H01M010-40

AB PURPOSE: To provide a nonaqueous electrolyte secondary battery with high capacity recovery capability and high charge/discharge cycle

performance by preventing the deposition of metallic lithium on the surface of an electrode by use of a negative electrode made mainly of mesophase graphite.

CONSTITUTION: Mesophase spherules are produced by heat melting treatment of coal pitch. The mesophase spherules are crushed, carbonized, and graphitized to produce mesophase graphite. Paste mainly comprising the mesophase graphite is applied to both surfaces of a foil core material of copper and the like and they are press molded to form a negative electrode. The negative electrode and a positive electrode are spirally wound through a separator to form an electrode group 4. From the electrode group 4, a positive lead 5 is connected to a sealing plate 2, and a negative lead 6 to the bottom of a battery case 1. The upper and lower parts of the electrode group 4 are covered with an insulating ring 7 respectively. They are housed in the battery case 1, and the battery case 1 is sealed with the sealing plate 2 through an insulating gasket 3. A secondary battery having high capacity, high energy density, and high charging performance at low temperature is provided.

COPYRIGHT: (C)1995,JPO

L35 ANSWER 35 OF 53 JAPIO (C) 2005 JPO on STN

AN 1995-153486 JAPIO

TI LITHIUM SECONDARY BATTERY

IN TSUKAMOTO HISASHI

PA JAPAN STORAGE BATTERY CO LTD

PI JP 07153486 A 19950616 Heisei

AI JP 1993-326273 (JP05326273 Heisei) 19931130

PRAI JP 1993-326273 19931130

SO PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined Applications, Vol. 1995

IC ICM H01M010-36

ICS H01M010-40

AB PURPOSE: To improve the high-rate discharge characteristic at a low temperature by using a lithium secondary battery added with gamma butyrolactone of a specific range % (volume %) to a mixture of ethylene carbonate and dimethyl carbonate at the specific volume ratio.

CONSTITUTION: The dielectric constant of an electrolyte added and mixed with gamma butyrolactone to a mixture of ethylene carbonate (EC) and dimethyl carbonate (DMC) at the volume ratio of 1:1 at -20°C is higher than that of the conventional mixture of EC+DMC+DEC (2:2:1). LiCoO<sub>2</sub> having the average grain size of 6 μm, acetylene black powder, and polyvinylidene fluoride are mixed, N-methyl pyrrolidone is added into a paste shape, and it is coated with an Al foil to form a positive electrode plate, for example. Artificial graphite and spherical graphite are mixed, polyvinylidene is mixed, N-methyl pyrrolidone is added, and it is applied to a copper foil to form a negative electrode plate. Both electrode plates are stored in a container, and the electrolyte is injected to obtain a battery having a high-rate discharge characteristic at a low temperature.

COPYRIGHT: (C)1995,JPO

L35 ANSWER 36 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1995:539256 HCAPLUS

DN 122:295263

ED Entered STN: 10 May 1995

TI Battery characteristics with various carbonaceous materials

AU Kuribayashi, Isao; Yokoyama, Mika; Yamashita, Masataka  
 CS A and T Battery Corporation, Imperial Tower 17F, 1-1 Uchisaiwai-cho,  
 1-chome, Chiyoda-ku, Tokyo, 100, Japan  
 SO Journal of Power Sources (1995), 54(1), 1-5  
 CODEN: JPSODZ; ISSN: 0378-7753  
 PB Elsevier  
 DT Journal  
 LA English  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy  
 Technology)  
 AB Various carbonaceous particles having a core-shell structure,  
 produced from **graphite** and **pseudo-graphite**  
 coated with **pitch**-blended phenol resin and green  
**mesophase-pitch** coated with phenol resin and  
 ground **graphite** powder, were investigated as active  
 materials for neg. **electrodes**. The particles with natural  
**graphite** cores, as well as those with pseudo-  
**graphite** cores, were stable both in 1M LiClO<sub>4</sub>-PC (propylene  
 carbonate):EC (ethylene carbonate) (50:50) electrolyte and in 1.5M  
 LiBF<sub>4</sub>-PC:EC:BL ( $\gamma$ -butyrolactone) (25:25:50) electrolyte, and  
 showed excellent elec. capacities. Those produced from green  
**mesophase-pitch** cores were of markedly deformed  
 configuration and showed large capacity gains with only small addns.  
 of **graphite** powder to the phenol coating material. Neg.  
**electrodes**, in which the particles contg. natural  
**graphites** cores were blended with petroleum coke in various  
 ratios, showed that it may be possible to produce cells with  
 specific discharge voltage curves, by varying the ratio of this  
 blend.  
 ST carbonaceous material anode battery  
 IT Carbonaceous materials  
 RL: DEV (Device component use); USES (Uses)  
 (battery characteristics with various carbonaceous materials)  
 IT Phenolic resins, uses  
 RL: DEV (Device component use); USES (Uses)  
 (carbonaceous particles coated with pitch-blended; battery  
 characteristics with various carbonaceous materials)  
 IT Anodes  
 (battery, battery characteristics with various carbonaceous  
 materials)  
 IT Coke  
 RL: DEV (Device component use); USES (Uses)  
 (petroleum, battery characteristics with various carbonaceous  
 materials)  
 IT 7782-42-5, Graphite, uses  
 RL: DEV (Device component use); USES (Uses)  
 (natural and artificial; battery characteristics with various  
 carbonaceous materials)  
 L35 ANSWER 37 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 1995:232637 HCAPLUS  
 DN 122:35101  
 ED Entered STN: 08 Dec 1994  
 TI Carbon fibers and natural graphite as negative electrodes for  
 lithium ion-type batteries  
 AU Yazami, R.; Zaghib, K.; Deschamps, M.  
 CS Laboratoire d'Ionique et d'Electrochimie du Solide de Grenoble (URA  
 CNRS 1213), ENSEEG, BP 75, Saint-Martin-d'Heres, 38402, Fr.  
 SO Journal of Power Sources (1994), 52(1), 55-60  
 CODEN: JPSODZ; ISSN: 0378-7753  
 PB Elsevier  
 DT Journal

LA English  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 Section cross-reference(s): 72  
 AB Carbon fibers (CFs), from different origins, and natural graphite are used as host lattices for lithium electrochem. intercalation and de-intercalation in org. liq. and solid polymer electrolytes, resp. In both systems, irreversible behavior occurred during the first cycle of which the origin is tentatively discussed. The reversible capacity of the mesophase CFs-based electrodes, which is related to the total relative amt. of lithium exchanged with the electrolyte during the charge/discharge operations, is found to increase with their crystallinity. Some thermodyn. data assocd. with the formation of the stage-1 graphite-lithium compd. in polymer electrolyte-based cells are detd.  
 ST lithium battery anode carbon fiber graphite  
 IT Carbon fibers, uses  
 RL: DEV (Device component use); USES (Uses)  
 (carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 IT Pitch  
 (isotropic, precursor for carbon fibers; carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 IT Batteries, secondary  
 (lithium/PEO-lithium perchlorate/graphite lithium and lithium/propylene carbonate-ethylene carbonate-dimethoxyethane-lithium tetrafluoroborate/carbon fibers)  
 IT Anodes  
 (battery, carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 IT Pitch  
 (mesophase, precursor for carbon fibers; carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 IT 12192-58-4 39448-96-9, Graphite lithium  
 RL: DEV (Device component use); USES (Uses)  
 (carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 IT 7782-42-5, Graphite, uses  
 RL: DEV (Device component use); USES (Uses)  
 (natural and pyrolytic; carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 IT 25014-41-9, Polyacrylonitrile  
 RL: NUU (Other use, unclassified); USES (Uses)  
 (precursor for carbon fibers; carbon fibers and natural graphite as neg. electrodes for lithium ion-type batteries)  
 L35 ANSWER 38 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 1994:58595 HCAPLUS  
 DN 120:58595  
 ED Entered STN: 05 Feb 1994  
 TI Secondary nonaqueous batteries and manufacture of anodes for the batteries  
 IN Ito, Zenichiro; Ozaki, Yoshuki; Morita, Teruyoshi; Yamaura, Junichi  
 PA Matsushita Electric Ind Co Ltd, Japan  
 SO Jpn. Kokai Tokkyo Koho, 10 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM H01M004-04  
 ICS H01M004-02; H01M004-58



CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05290833	A2	19931105	JP 1992-90534	19920410
	JP 3191394	B2	20010723		
	US 5344724	A	19940906	US 1993-42794	19930406

PRAI JP 1992-90534 A 19920410

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 05290833	ICM	H01M004-04
	ICS	H01M004-02; H01M004-58
	IPCI	H01M0004-04 [ICM,5]; H01M0004-02 [ICS,5]; H01M0004-58 [ICS,5]
US 5344724	IPCI	H01M0004-58 [ICM,5]; H01M0006-14 [ICS,5]
	NCL	429/094.000; 429/231.300; 429/231.800
	ECLA	H01M004/58E2; H01M010/40

AB Mesophase granules sepd. from the melted pitch is carbonized and graphitized at 2500-2900° to form graphite granules, mixed with a binder, applied on collectors, and pressed to obtain the title anodes having 25-40% porosity. The batteries use Li-contg. metal oxide-based cathodes and graphite anodes, where the graphite has interplanar spacing d002 3.365-3.390 Å, unit cell length Lc 200-650 Å, and I1360/I1580 = 0.20.apprx.0.40 ( I1360 and I1580 are the intensities of peaks at 1360 and 1580/cm, resp., on the Ar laser Reman spectrum of the graphite).

ST lithium battery graphite anode

IT Carbonaceous materials

RL: USES (Uses)  
(graphitized, specifications and manuf. of, for lithium battery anodes)

IT Anodes  
(battery, lithium, graphitized carbonaceous materials for, specifications and prepn. of)

IT 7782-42-5, Graphite, uses  
RL: DEV (Device component use); USES (Uses)  
(for anodes, specifications and manuf. of, for lithium batteries)

L35 ANSWER 39 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1993:630107 HCAPLUS

DN 119:230107

ED Entered STN: 27 Nov 1993

TI Secondary organic-electrolyte batteries

IN Yasunami, Shoichiro; Kagawa, Okimasa; Maekawa, Yukio

PA Fuji Photo Film Co Ltd, Japan

SO Jpn. Kokai Tokkyo Koho, 12 pp.  
CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM H01M004-58

ICS H01M004-02; H01M010-40

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 05174820	A2	19930713	JP 1991-338959	19911220
	JP 3239302	B2	20011217		
PRAI	JP 1991-338959		19911220		

## CLASS

	PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
	JP 05174820	ICM	H01M004-58
		ICS	H01M004-02; H01M010-40
		IPCI	H01M0004-58 [ICM,5]; H01M0004-02 [ICS,5]; H01M0010-40 [ICS,5]
		ECLA	H01M004/48B2; H01M004/52B2; H01M004/58E; H01M004/58E2
AB	The batteries have Li-contg. transition metal chalcogenide cathodes and carbonaceous anodes, where the anodes are composed of fine low-graphitization C powder having unit cell length Lc 8-150Å, interplanar spacing d002 3.42-3.65Å, real d. 1.60-2.20 g/cm <sup>3</sup> and/or fine carbon fibers. The C powder may be carbonized polyacrylonitrile, coal-based coke, mesophase pitch, or carbon black; and the carbon fibers may be fibrous graphite. The chalcogenide may be LiaCobVcOd (a=0.1-1.1, b=0.15-0.9, c=1-b, d=2-2.5) or LiaCobNicOd.		
ST	lithium battery carbonaceous anode; transition metal lithium oxide cathode		
IT	Carbon black, uses Coke RL: USES (Uses) (anodes contg., lithium-intercalating, for batteries)		
IT	Batteries, secondary (lithium-intercalating carbon/lithium transition metal chalcogenide)		
IT	Carbon fibers, uses RL: USES (Uses) (graphite, anodes contg., lithium-intercalating, for batteries)		
IT	Pitch (mesophase, anodes contg., lithium-intercalating, for batteries)		
IT	9003-54-7D, pyrolyzed 25014-41-9D, Polyacrylonitrile, pyrolyzed RL: USES (Uses) (anodes contg., lithium-intercalating, for batteries)		
IT	7439-93-2, Lithium, uses RL: USES (Uses) (anodes from carbonaceous materials intercalated with, for batteries)		
IT	7440-44-0 7782-42-5 RL: USES (Uses) (carbon fibers, graphite, anodes contg., lithium-intercalating, for batteries)		
IT	147521-85-5, Cobalt lithium vanadium oxide (Co <sub>0.5</sub> Li <sub>0.5</sub> V <sub>0.5</sub> O <sub>2.5</sub> ) 151028-39-6, Cobalt lithium nickel oxide (Co <sub>0.5</sub> Li <sub>0.5</sub> Ni <sub>0.5</sub> O <sub>2.5</sub> ) RL: DEV (Device component use); USES (Uses) (cathodes, for batteries)		

L35 ANSWER 40 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1994:139205 HCAPLUS

DN 120:139205

ED Entered STN: 19 Mar 1994

TI Lithium secondary battery and method of manufacturing carbonaceous anode materials

IN Takami, Norio; Satoh, Asako; Ohsaki, Takahisa  
 PA Toshiba Corp., Japan  
 SO Eur. Pat. Appl., 52 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA English  
 IC ICM H01M004-58  
 ICS H01M010-40  
 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 573266	A1	19931208	EP 1993-304258	19930601
	EP 573266	B1	19991208		
	R: DE, FR, GB				
	JP 06168725	A2	19940614	JP 1993-56194	19930316
	JP 3241850	B2	20011225		
	JP 06275321	A2	19940930	JP 1993-58565	19930318
	JP 3162531	B2	20010508		
	US 5340670	A	19940823	US 1993-69424	19930601
	KR 134638	B1	19980515	KR 1993-9983	19930601
PRAI	JP 1992-140247	A	19920601		
	JP 1992-261418	A	19920930		
	JP 1993-56194	A	19930316		
	JP 1993-58565	A	19930318		

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
EP 573266	ICM	H01M004-58
	ICS	H01M010-40
	IPCI	H01M0004-58 [ICM,5]; H01M0010-40 [ICS,5]
	ECLA	H01M004/58E2; H01M010/40
JP 06168725	IPCI	H01M0004-58 [ICM,5]; H01M0004-02 [ICS,5]; H01M0010-40 [ICS,5]
JP 06275321	IPCI	H01M0010-40 [ICM,5]; H01M0004-02 [ICS,5]; H01M0004-58 [ICS,5]
US 5340670	IPCI	H01M0010-40 [ICM,5]
	NCL	429/331.000; 423/445.000R; 423/448.000; 429/231.500; 429/231.800; 429/330.000; 429/332.000
KR 134638	IPCI	H01M0004-02 [ICM,6]

AB The lithium secondary battery includes a **cathode** housed in a case, an anode housed in the case and contg. a carbonaceous material with exothermic peak at  $\geq 700^\circ$  when measured by a differential thermal anal. and an intensity ratio P101/P100 of a (101) diffraction peak P101 to a (100) diffraction peak P100 of a **graphite** structure, obtained by x-ray diffraction anal., of 0.7 to 2.2, and absorbs lithium ions, a separator housed in the case so as to be arranged between the pos. and neg. **electrodes**, and a nonaq. electrolyte contained in the case. The battery has high capacity and excellent charge-discharge efficiency, cycle life,

flatness of a discharge voltage, and rapid charge-discharge cycle characteristics.

ST lithium battery carbon anode

IT Rubber, butadiene-styrene, uses  
RL: PREP (Preparation)  
(in carbon anode prepn. for nonaq. secondary lithium batteries)

IT Carbon black, uses  
RL: USES (Uses)  
(in nonaq. secondary lithium batteries with carbon anode)

IT Batteries, secondary  
(lithium, nonaq., with carbon anode)

IT Pitch  
(mesophase, in carbon anode prepn. for nonaq. secondary lithium batteries)

IT 9004-32-4, Cmc  
RL: USES (Uses)  
(in carbon anode prepn. for nonaq. secondary lithium batteries)

IT 75-05-8, Acetonitrile, uses 75-52-5, Nitromethane, uses 79-20-9, Methyl acetate 96-48-0,  $\gamma$ -Butyrolactone 96-49-1, Ethylene carbonate 98-95-3, Nitrobenzene, uses 105-58-8, Diethyl carbonate 107-21-1, Ethylene glycol, uses 108-32-7, Propylene carbonate 108-88-3, Toluene, uses 141-78-6, Ethyl acetate, uses 616-38-6, Dimethyl carbonate 1330-20-7, Xylene, uses 7429-90-5, Aluminum, uses 7440-02-0, Nickel, uses 7440-50-8, Copper, uses 7782-50-5, Chlorine, uses 7791-03-9, Lithium perchlorate 9002-84-0, Ptfе 9010-79-1, Ethylene-propylene copolymer 12597-68-1, Stainless steel, occurrence 14283-07-9, Lithium borofluoride 21324-40-3 29935-3 5-1 33454-82-9 39300-70-4, Lithium nickel oxide 39457-42-6, Lithium manganese oxide 52627-24-4, Lithium cobalt oxide 90076-65-6, Lithium bistrifluoromethylsulfonylimide  
RL: USES (Uses)  
(in nonaq. secondary lithium batteries with carbon anode)

IT 9003-55-8  
RL: USES (Uses)  
(rubber, in carbon anode prepn. for nonaq. secondary lithium batteries)

L35 ANSWER 41 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1993:201212 HCAPLUS

DN 118:201212

ED Entered STN: 14 May 1993

TI Manufacture of artificial graphite electrodes for electrolytic steel manufacture

IN Kanbe, Masazumi; Noda, Tomoyoshi; Miwa, Shigeru

PA Nippon Steel Chemical Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.  
CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C25C007-02  
ICS C01B031-04; C04B035-54

CC 72-2 (Electrochemistry)  
Section cross-reference(s): 55

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 04285189	A2	19921009	JP 1991-70316	19910312
PRAI JP 1991-70316		19910312		

CLASS  
PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES  
-----

JP 04285189 ICM C25C007-02  
ICS C01B031-04; C04B035-54  
IPCI C25C0007-02 [ICM,5]; C01B0031-04 [ICS,5];  
C04B0035-54 [ICS,5]

AB In the manuf. of artificial **graphite electrodes**  
using needle coke and binder **pitch** as starting materials,  
 $\beta$  resin conc. is added when needle coke and binder  
**pitch** are kneaded and heated or before the kneading and  
heating it is added to binder **pitch** and/or needle coke,  
thereby the content of  $\beta$  resin in the entire blend is adjusted  
to 6-10 wt.%. The  $\beta$  resin conc. is obtained from the extn. of  
coal tar or coal tar **pitch**es by arom. hydrocarbon solvents  
and contains ash  $\leq 0.1$  wt.% and quinoline-sol. but  
toluene-insol matter  $\geq 50$  wt.% followed by pulverization. The  
content of  $\beta$  resin is a kneaded mixt. of starting materials can  
be adjusted easily and it is unnecessary to carry out excessive  
thermal modification which can bring about softening point rising or  
formation of **mesophase** by the increase of  $\beta$  resin in  
binder **pitch** so that workability becomes better and  
artificial **graphite electrodes** having improved  
properties can be obtained just by adjusting the amt. of  $\beta$   
resin conc. to be added.

ST graphite electrode beta resin adjustment; coke needle binder pitch  
graphite electrode; steel electrochem prepn graphite electrode

IT Electrodes  
(graphite, artificial, prepn. of, from needle coke and binder  
pitch for steel making)

IT Polymers, uses  
RL: USES (Uses)  
( $\beta$ -, in artificial graphite electrodes manuf., phys.  
properties of electrodes in relation to)

IT 7782-42-5P, Graphite, uses  
RL: PREP (Preparation)  
(electrodes, artificial, prepn. of, from needle coke and binder  
pitch)

IT 12597-69-2P, Steel, preparation  
RL: PREP (Preparation)  
(manuf. of, electrolytic, artificial graphite electrode for)

L35 ANSWER 42 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
AN 1991:252629 HCAPLUS  
DN 114:252629  
ED Entered STN: 28 Jun 1991  
TI Manufacture of graphite **electrode nipples**  
IN Griffin, Robert R.; Roussel, Keith M.; Fu, Ta Wei  
PA Conoco, Inc., USA  
SO U.S., 5 pp. cont.-in-part of U.S. Ser. No. 210,689, abandoned.  
CODEN: USXXAM  
DT Patent  
LA English  
IC ICM C01B031-00  
INCL 264029500  
CC 57-8 (Ceramics)  
Section cross-reference(s): 55  
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI US 4998709	A	19910312	US 1990-513468	

JP 04230994 A2 19920819 JP 1991-59588 199004  
19  
JP 3351540 B2 20021125 199102  
PRAI US 1988-210689 B2 19880623 28  
US 1990-513468 A 19900419

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
US 4998709	ICM	C01B031-00
	INCL	264029500
	IPCI	C01B0031-00 [ICM,5]
	NCL	264/029.500; 029/825.000; 264/029.700; 264/105.000; 264/108.000; 373/092.000; 403/DIG.005; 428/408.000
JP 04230994	IPCI	H05B0007-14 [ICM,5]; C25B0009-02 [ICA,5]
AB		<b>Graphite</b> nipples suitable for connecting <b>graphite electrodes</b> used in steelmaking are prep'd. by adding 8-20 wt.% C fibers produced from spun <b>mesophase pitch</b> to 45-65 wt.% premium coke and 22-28 wt.% binder to form an extrusion blend, extruding the extrusion blend to form an elongated C nipple, and subjecting the C nipple to 760-982° for 2-5 wk then to .gtorsim.2704° for 5-14 days. The graphitized C fibers of the <b>graphite</b> nipples have tensile strength (200-300) + 103 psi and Young's modulus of elasticity (55-120) + 106 psi.
ST		<b>graphite electrode joint</b> prepn steelmaking; <b>mesophase pitch graphite electrode joint</b> ; coke <b>graphite electrode joint</b> ; carbon fiber <b>graphite electrode joint</b>
IT		Joints, mechanical ( <b>electrodes, graphite nipples</b> for, prepn. of, <b>mesophase pitch-based carbon fibers</b> in, for high strength)
IT		Coke RL: SPN (Synthetic preparation); PREP (Preparation) (in <b>graphite electrode joint</b> prepn. from <b>mesophase pitch-based carbon fibers</b> )
IT		<b>Pitch</b> (coal-tar, in <b>graphite electrode joint</b> prepn. from <b>mesophase pitch-based carbon fibers</b> )
IT		<b>Electrodes</b> (furnace, <b>graphite nipples</b> for, manuf. of, for steelmaking)
IT		<b>Pitch</b> ( <b>mesophase, carbon fibers</b> from, manuf. of, in <b>graphite electrode joint</b> prepn.)
IT		Carbon fibers, uses and miscellaneous RL: SPN (Synthetic preparation); PREP (Preparation) ( <b>pitch-based, mesophase, in graphite electrode joint</b> prepn.)
IT		7440-44-0 7782-42-5 RL: USES (Uses) (carbon fibers, <b>pitch-based, mesophase, in graphite electrode joint</b> prepn.)
IT		1332-37-2, Iron oxide, uses and miscellaneous RL: USES (Uses) (in <b>graphite electrode joint</b> prepn. from <b>mesophase pitch-based carbon fibers</b> )

L35 ANSWER 43 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1990:636597 HCAPLUS  
 DN 113:236597  
 ED Entered STN: 22 Dec 1990  
 TI Manufacture of graphite products with high density and strength, and  
 graphite electrodes for electric-discharge machining  
 IN Fukuda, Noriyoshi; Hagiwara, Kiyoshi; Torii, Takayuki; Nagayama,  
 Katsuhiko  
 PA Kawasaki Steel Corp., Japan  
 SO Jpn. Kokai Tokkyo Koho, 11 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM C04B035-54  
 ICS B23H001-08; C10C003-00  
 CC 57-8 (Ceramics)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 02088464	A2	19900328	JP 1988-163626	19880630
PRAI	JP 1988-33750	A1	19880216		
	JP 1988-136590	A1	19880602		

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 02088464	ICM	C04B035-54
	ICS	B23H001-08; C10C003-00
	IPCI	C04B0035-54 [ICM,5]; B23H0001-08 [ICS,5]; C10C0003-00 [ICS,5]

AB The title **graphite** products (or **electrodes**) are  
 manufd. by mixing **mesophase** C beads with  $\leq 40$  wt.%  
 mixt. contg. 100 wt. parts **graphite** and/or coke and 30-120  
 wt. parts **pitch** contg. from 40 to  $< 95$  wt.% C6H6 insols.,  
 molding, firing, and graphitizing.  
 ST **mesophase** carbon **pitch** coke **graphite**  
 product; **electrode** **graphite** elec discharge  
 machining  
 IT Pitch  
 (graphite products from mesophase carbon and coke and, with high  
 d. and strength)  
 IT Coke  
 RL: USES (Uses)  
 (graphite products from mesophase carbon and pitch and, with high  
 d. and strength)  
 IT Electrodes  
 (graphite, manuf. of, for elec.-discharge machining)  
 IT Machining  
 (elec.-discharge, **graphite electrodes** for,  
 manuf. of high-d. and -strength, from **mesophase** carbon  
 and coke and **pitch**)  
 IT 7440-44-0, Carbon, uses and miscellaneous  
 RL: USES (Uses)  
 (mesophase, graphite products from coke and pitch and, with high  
 d. and strength)  
 IT 7782-42-5, Graphite, uses and miscellaneous  
 RL: USES (Uses)  
 (products, with high d. and strength, from mesophase carbon and  
 coke and pitch)

L35 ANSWER 44 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1989:234523 HCAPLUS  
 DN 110:234523  
 ED Entered STN: 25 Jun 1989  
 TI **Mesophase pitch for graphite electrode production**  
 IN Matsubara, Kenji; Okuyama, Yasuo; Ueno, Ichiro  
 PA Nippon Kokan K. K., Japan  
 SO Jpn. Kokai Tokkyo Koho, 3 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM C10C003-04  
 ICS C10C003-10  
 ICA C04B041-82; C10C003-02  
 CC 51-19 (Fossil Fuels, Derivatives, and Related Products)  
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63278995	A2	19881116	JP 1987-114009	19870511
JP 04038790	B4	19920625		
PRAI JP 1987-114009		19870511		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 63278995	ICM	C10C003-04
	ICS	C10C003-10
	ICA	C04B041-82; C10C003-02
	IPCI	C10C0003-04 [ICM,4]; C10C0003-10 [ICS,4]; C04B0041-82 [ICA,4]; C10C0003-02 [ICA,4]

AB The title pitch is manufd. by hydrogenating petroleum pitch or coal-tar pitch, mixing 5-40% light oil with the hydrogenated pitch and heating the mixt. under inert atm. environment to obtain a mesophase pitch product. The light oil has an av. mol. wt. of 100-500 and contains mainly 2-6-ring arom. components. Thus, a coal-tar pitch was catalytically hydrogenated, mixed with 25% light oil (b. 540°), and heated at 400° under N for .apprx.2 h to obtain a mesophase pitch with viscosity 0.25 P.  
 ST **pitch mesophase graphite electrode prodn; coal tar pitch hydrogenation mesophase**  
 IT **Electrodes**  
 (graphite, prodn. of, mesophase pitch for)  
 IT **Pitch**  
 (mesophase, manuf. of, from coal-tar pitch, for graphite electrode prodn.)

L35 ANSWER 45 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 1989:176541 HCAPLUS  
 DN 110:176541  
 ED Entered STN: 12 May 1989  
 TI Coal-tar pitch for graphite electrode production  
 IN Sato, Maki; Matsui, Yoshiaki; Yamada, Masahiro; Fujimoto, Kenichi  
 PA Nippon Steel Corp., Japan; Nippon Steel Chemical Co., Ltd.  
 SO Jpn. Kokai Tokkyo Koho, 8 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
 IC ICM C10C003-04



ICS C01B031-00; C10C003-10  
 CC 51-19 (Fossil Fuels, Derivatives, and Related Products)  
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63256690	A2	19881024	JP 1987-90673	19870415

JP 06072224 B4 19940914  
 PRAI JP 1987-90673 19870415

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 63256690	ICM	C10C003-04
	ICS	C01B031-00; C10C003-10
	IPCI	C10C0003-04 [ICM,4]; C01B0031-00 [ICS,4]; C10C0003-10 [ICS,4]

AB The title pitch is prepd. by (a) contacting the coal-tar pitch with a hydrogenation catalyst at 300-500° to obtain a hydrogenated pitch having an H absorption amt. 5 g/kg pitch feed and the denitrogenation efficiency ≤80 wt.%; and (b) heat-soaking the hydrogenated pitch at 350-450° under N for 0.5-10 h, preferably 4-6 h to obtain a mesophase pitch product. Thus, a coal-tar pitch (softening point 31.6°, contg. MePh-insols. 6.8, quinoline-insols. trace, fixed C 31.8, and N 1.46 wt.%) was contacted with a Ni-Mo/Al<sub>2</sub>O<sub>3</sub> catalyst at 398°, 150 kg/cm<sup>2</sup> and 1.98 h-l liq. space velocity to obtain a soft hydrogenated pitch, which was then heated under N in an autoclave at 380° for 5 h to obtain a mesophase pitch having softening point 90.6°, contg. MePh-insols. 29.6, quinoline-insols. 9.8 and fixed C 57.1 wt.%.

ST coal tar pitch mesophase manuf; graphite  
 electrode mesophase pitch hydrogenation;  
 nickel molybdenum catalyst pitch hydrogenation

IT Pitch  
 (mesophase, manuf. of, from coal-tar pitch,  
 by hydrogenation and heat-soaking, for graphite  
 electrode prodn.)

IT 7439-98-7, Molybdenum, uses and miscellaneous 7440-02-0, Nickel,  
 uses and miscellaneous  
 RL: CAT (Catalyst use); USES (Uses)  
 (catalyst contg., in hydrogenation of coal-tar pitch)

L35 ANSWER 46 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1989:176540 HCAPLUS

DN 110:176540

ED Entered STN: 12 May 1989

TI Coal-tar pitch for graphite electrode production

IN Matsui, Yoshiaki; Sato, Maki; Yamada, Masahiro; Fujimoto, Kenichi

PA Nippon Steel Corp., Japan; Nippon Steel Chemical Co., Ltd.

SO Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C10C003-02

ICS C01B031-00; C10C003-10

CC 51-19 (Fossil Fuels, Derivatives, and Related Products)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 63256689 A2 19881024 JP 1987-90674 198704  
15

JP 06072226 B4 19940914  
PRAI JP 1987-90674 19870415

CLASS  
PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES  
-----  
JP 63256689 ICM C10C003-02  
ICS C01B031-00; C10C003-10  
IPCI C10C0003-02 [ICM,4]; C01B0031-00 [ICS,4];  
C10C0003-10 [ICS,4]

AB The title pitch is manufd. by blending the pitch feed with  
≥10 wt.% alcs., heat-soaking the mixt. at  
≥250°, and distg. the reaction product to obtain a  
mesophase pitch with lower softening point. The heat-soaking is  
preferably carried out at 330-450° for 1-10 h. Thus, 100 wt.  
parts coal-tar pitch and 45 wt. parts 2-propanol were fed into an  
autoclave, heated under air, and then N atm. at 360° and 150  
kg/cm2 for 5 h to obtain a mesophase pitch with softening point  
89.6° vs. 91.2° for a conventional method.

ST coal tar pitch mesophase electrode;  
propanol pitch mesophase graphite  
electrode

IT Pitch  
(mesophase, manuf. of, from coal-tar pitch,  
by heat-soaking in presence of alcs., for graphite  
electrode prodn.)

IT 67-63-0, 2-Propanol, uses and miscellaneous 107-21-1, Ethylene  
glycol, uses and miscellaneous 141-43-5, Ethanolamine, uses and  
miscellaneous

RL: USES (Uses)  
(coal-tar pitch heat soaking with, in prepn. of  
mesophase pitch, for graphite  
electrode prodn.)

L35 ANSWER 47 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1988:476486 HCAPLUS

DN 109:76486

ED Entered STN: 02 Sep 1988

TI Method for manufacture of carbonaceous feedstock powders used in the  
production of high-density graphite electrodes

IN Wakasa, Tsutomu; Ono, Taku; Yamamoto, Tsuneo; Wakabayashi, Takashi

PA Nippon Denkyoku K. K., Japan

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C01B031-02

CC 51-19 (Fossil Fuels, Derivatives, and Related Products)  
Section cross-reference(s): 57

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63089413	A2	19880420	JP 1986-233821	198610 01

PRAI JP 1986-233821 19861001

CLASS  
PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES  
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JP 63089413 ICM C01B031-02

IPCI C01B0031-02 [ICM,4]

AB The title method comprises (a) bubbling molten coal-tar pitch with N gas at 300-420° to remove volatile org. components in a 1st reforming stage; (b) heating the reformed pitch at 400-450°, and then atomizing through a spray nozzle under N atm. to obtain a reformed pitch powder (av. diam. 1-100 µm); (c) contacting the reformed pitch powder with hot air in a fluidized bed at 200-300° to form a high m.p. pitch powder in a 2nd reforming stage; and (d) heating the high m.p. pitch powder under N at 480-550° for 0.5-3 h to obtain mesophase pitch product useful in the prodn. of high-d. graphite electrodes.

ST coal tar pitch mesophase manuf; graphite electrode mesophase pitch

IT Pitch  
(mesophase, manuf. of, from coal-tar pitch, for graphite electrode prodn.)

IT 7782-42-5P, uses and miscellaneous  
RL: PREP (Preparation)  
(electrodes, prodn. of, mesophase pitch prepn. in)

L35 ANSWER 48 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1986:556186 HCAPLUS

DN 105:156186

ED Entered STN: 01 Nov 1986

TI Graphite-base plates

IN Takahashi, Kunimasa; Kameda, Takashi; Shibatani, Haruo

PA Mitsubishi Petrochemical Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C04B035-54

ICS C01B031-04

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

Section cross-reference(s): 51, 57

FAN.CNT 4

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	---	-----	-----	
PI	JP 61077667	A2	19860421	JP 1984-199737	198409 25
	US 4929404	A	19900529	US 1988-196760	198805 17
PRAI	JP 1984-199737	A	19840925		
	JP 1984-255270	A	19841203		
	JP 1985-63329	A	19850329		
	JP 1985-87264	A	19850423		
	US 1985-779590	B1	19850924		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
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JP 61077667	ICM	C04B035-54
	ICS	C01B031-04
	IPCI	C04B0035-54 [ICM,4]; C01B0031-04 [ICS,4]
US 4929404	IPCI	C10C0003-00 [ICM,5]; B29C0043-52 [ICS,5]

NCL 264/029.500; 208/044.000; 264/029.100;  
264/029.700; 264/101.000; 264/DIG.020

- AB The mold-formed plates have elec. resistivity  $\leq 5.0$  m $\Omega$ -cm and bending strength  $\geq 200$  kg/cm<sup>2</sup>, and they show vol. and wt. change  $\leq 3\%$  when heated at 1000° to be carbonized. Powder materials (max. m.p. 400°, carbonizing yield  $\geq 70\%$  at 1000°) of **graphite** and **pitch** contg.  $\leq 70\%$  quinoline-insol. component and  $\geq 40\%$  **mesophase** are compacted and fired in inert atm. at  $\geq 700^\circ$ . The dimensionally stable plates are useful as **electrode** substrates and separators of fuel cells. Thus, 630 g tar (b.p.  $\geq 170^\circ$ , thermal decompn. residue of naphtha) was treated with H at 120 kg/cm<sup>2</sup> and 460° in the presence of cracking catalysts (Al<sub>2</sub>O<sub>3</sub>-SiO<sub>2</sub>), filtered, and distd. at  $\leq 490^\circ$  to give reformed **pitch** in 25%. The **pitch** (10 g) was placed in a reaction vessel, treated with 1,2,3,4-tetrahydroquinoline (I) under an Ar flow for 10 min, immersed in a molten salt bath at 485° while I and Ar were supplied to the liq. **pitch** to give **pitch** (in 52% yield) contg. 100 **mesophase** and 45% quinoline-insol. component. The **mesophase**-contg. **pitch** (1.34 g) was mixed with 8.0 g CPB flake **graphite**, ground, and 1.70 g of the mixt. was packed into a metal mold and precompacted at 1.5 ton/cm<sup>2</sup> and 340°. The green compact was held at 1000° under an Ar flow for 30 min to give 1.67 g **graphite**-base plate having bulk d. 2.1 g/mL, vol. shrinkage 0.6%, wt. decrease 1.7% (based on the green compact), elec. resistivity 1.2 m $\Omega$ -cm, and bending strength 550 kg/cm<sup>2</sup>.
- ST **graphite pitch fuel cell electrode; separator fuel cell graphite pitch; resistance graphite pitch plate**
- IT **Electric resistance**  
(of graphite plates, for fuel-cell electrodes and separators)
- IT **Fuel cells**  
(separators graphite plates for, pitch-bonded stable and high-d.)
- IT **Electrodes**  
(fuel-cell, graphite plates for, pitch-bonded stable, and high d.)
- IT **Pitch**  
(petroleum, **mesophase**, binders, for **graphite** plates, for fuel-cell **electrodes** and separators)
- IT **7782-42-5P, preparation**  
RL: PREP (Preparation)  
(manuf. of plates contg., with **mesophase**-contg. **pitch** binder, for fuel-cell **electrodes** and separators)

L35 ANSWER 49 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1985:426257 HCAPLUS

DN 103:26257

ED Entered STN: 27 Jul 1985

TI Formed carbon as **electrodes**

PA Koa Oil Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM C01B031-04

CC 57-8 (Ceramics)

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 60016806 A2 19850128 JP 1983-122565 198307  
06

JP 01023405 B4 19890502  
PRAI JP 1983-122565 19830706

## CLASS

PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES

JP 60016806 ICM C01B031-04  
IPCI C01B0031-04 [ICM,4]

AB To graphitize effectively, a bulk mesophase material (A) (i.e. mesocarbon microbeads prep'd. by heating heavy oil at 400-500° and sepg.) is mixed at ≤20% with coal or petroleum pitch as binder (or with acicular coke or their mixt.). Thus, coal-tar pitch (softening temp. 109°, fixed C 56, quinoline-insolubles 4.2%) was mixed with 3 or 10% A -60 mesh, kneaded with com. needle petroleum coke (8-20, 20-35, and -100 mesh 30, 10, and 60%, resp.) in a 28:100 ratio at 160° for 1 h, extruded at 130°, heated at 200°/h to and at 1000° for 1 h, and then to 2800° to graphitize. The d. was 1.51 or 1.54, Young's modulus 730 or 890, bending strength 100 or 120 kg/mm<sup>2</sup>, elec. resistivity (+ 10<sup>-3</sup> Ω-cm) 0.9 each, thermal expansion coeff. 0.46 or 0.49 + 10<sup>-6</sup>/degree, and the calcd. thermal shock resistance 920 or 860 cal/cm-s, compared to 1.48, 690, 90, 1.0, 0.45, and 860, resp., with A.

ST carbon mesophase material graphitization electrode

IT Electrodes  
(graphite, mesophase microbead additives in manuf. of)

IT Graphitization  
(of carbon, mesophase microbead additives in, for electrodes)

IT Pitch  
(coal-tar, graphite electrode manuf. from, mesophase microbead additives in)

IT Spheres  
(micro-, mesophase carbon, in graphite electrode manuf.)

IT Coke  
RL: USES (Uses)  
(petroleum, graphite electrode manuf. from, mesophase microbead additives in)

IT 7440-44-0, properties  
RL: RCT (Reactant); RACT (Reactant or reagent)  
(graphitization of, mesophase microbead additive in, for electrodes)

L35 ANSWER 50 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1979:576486 HCAPLUS

DN 91:176486

ED Entered STN: 12 May 1984

TI Elastomer piezoresistors

IN Kanamori, Katsuhiko

PA Yokohama Rubber Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC C08K007-00; H01B001-04

CC 38-9 (Elastomers, Including Natural Rubber)

Section cross-reference(s): 76

FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	----- JP 54080350	A2	19790627	JP 1977-148045	197712 12
	JP 56009187	B4	19810227		
	GB 1561189	A	19800213	GB 1977-53596	197712 22
	CA 1096161	A1	19810224	CA 1977-293761	197712 22
	NL 7714355	A	19780627	NL 1977-14355	197712 23
	NL 190556	B	19931116		
	NL 190556	C	19940418		
	DE 2757870	A1	19780629	DE 1977-2757870	197712 23
	DE 2757870	C2	19870813		
	FR 2375698	A1	19780721	FR 1977-39231	197712 26
	FR 2375698	B1	19830429		
	US 4273682	A	19810616	US 1979-83367	197910 10
PRAI	JP 1976-155107	A	19761224		
	JP 1977-148045	A	19771212		
	US 1977-862855	A1	19771212		

## CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 54080350	IC	C08K0007-00; H01B001-04
	IPCI	C08K0007-00; H01B0001-04
GB 1561189	IPCI	C08K0007-00
CA 1096161	IPCI	H01B0001-00
NL 7714355	IPCI	C08K0009-00; C08K0003-04; H01C0010-12; H01B0017-64
DE 2757870	IPCI	C08K0007-18
FR 2375698	IPCI	H01C0010-10; B32B0019-02; C08K0003-04; C08K0007-00; C08L0027-06; C08L0083-04
US 4273682	IPCI	H01B0001-06
	NCL	252/511.000; 338/114.000

AB Rubbery polymers contg. powd. **synthetic graphite** having degree of roundness (Wadell) >0.5 are useful as piezoresistors. Thus, a compn. of KE 640-U (silicone rubber) 100, crushed and ground **synthetic graphite** (60-105  $\mu$ , degree of roundness 0.72) 70, and peroxide 3.4 parts was rolled and vulcanized to give a 0.5-mm sheet. When the sheet was tested for piezoresistivity between a flat **electrode** and a 3-mm-diam. rod **electrode** (max. pressure 1 kg), there was no noise for >106 cycles, compared with <103 cycles for a similar compn. contg. crushed **synthetic graphite** of degree of roundness 0.25.

ST silicone rubber graphite composite; piezoresistance silicone rubber composite; **roundness degree graphite**

IT Rubber, silicone, uses and miscellaneous  
RL: USES (Uses)  
(piezorsistors, contg. powd. graphite)

IT Piezoresistors  
(powd. graphite-contg. rubber)

IT 7782-42-5, uses and miscellaneous  
 RL: USES (Uses)  
 (piezoresistors contg., rubber-based)  
 IT 9002-86-2  
 RL: USES (Uses)  
 (piezoresistors, contg. powd. graphite)

L35 ANSWER 51 OF 53 HCAPLUS COPYRIGHT 2005 ACS on STN  
 AN 1976:484759 HCAPLUS  
 DN 85:84759  
 ED Entered STN: 12 May 1984  
 TI Improved graphite articles with low thermal expansion coefficients  
 IN Singer, Leonard S.; Lewis, Irwin Charles  
 PA Union Carbide Corp., USA  
 SO Ger. Offen., 23 pp.  
 CODEN: GWXXBX  
 DT Patent  
 LA German  
 IC C04B; C25B  
 CC 72-7 (Electrochemistry)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	---	-----	-----	
PI	DE 2542953	A1	19760408	DE 1975-2542953	197509 26
	DE 2542953	B2	19790104		
	DE 2542953	C3	19790830		
	CA 1060161	A1	19790814	CA 1975-234616	197508 29
	AU 7585172	A1	19770331	AU 1975-85172	197509 25
	BE 833903	A1	19760326	BE 1975-160447	197509 26
	DK 7504337	A	19760328	DK 1975-4337	197509 26
	SE 7510818	A	19760329	SE 1975-10818	197509 26
	SE 415094	B	19800908		
	SE 415094	C	19810108		
	NL 7511384	A	19760330	NL 1975-11384	197509 26
	NO 7503281	A	19760330	NO 1975-3281	197509 26
	NO 140716	C	19791024		
	NO 140716	B	19790716		
	FR 2286107	A1	19760423	FR 1975-29677	197509 26
	FR 2286107	B1	19800725		
	JP 51063809	A2	19760602	JP 1975-116292	197509 26
	JP 54026235	B4	19790903		

ZA 7506130	A	19760929	ZA 1975-6130	197509 26
ES 441286	A1	19770401	ES 1975-441286	197509 26
CH 605466	A	19780929	CH 1975-12516	197509 26
GB 1526809	A	19781004	GB 1975-39441	197509 26
AT 7507375	A	19830615	AT 1975-7375	197509 26

AT 373566	B	19840210
PRAI US 1974-510039	A	19740927

CLASS PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES

DE 2542953	IC	C04B; C25B
	IPCI	C04B0035-54; C25B0011-12
CA 1060161	IPCI	C01B0031-00
AU 7585172	IPCI	C09C0001-46
BE 833903	IPCI	C04B
DK 7504337	IPCI	C25B
SE 7510818	IPCI	C01B0031-04
NL 7511384	IPCI	C04B0035-54; C01B0031-04; C25B0011-12
NO 7503281	IPCI	C04B
FR 2286107	IPCI	C01B0031-04; H05B0031-08
JP 51063809	IPCI	C04B0035-64; C01B0031-04
ZA 7506130	IPCI	C10B
ES 441286	IPCI	C10B; B29F
CH 605466	IPCI	C04B0035-54
GB 1526809	IPCI	C01B0031-04
AT 7507375	IPCI	C01B0031-04

AB The title materials are applicable to the fabrication of electrodes for high temp. operation. Thus, from a com. petroleum pitch was made a pitch with a mesophase content of 57%, a d. of 1.24 and a softening point of 120°. The chem. anal. was C 93.3, H 5.63, S 1.0 and ash 0.15%. (The quinoline-insol. content was 0.5%). The mesophase pitch was made by heating pitch for 15 hr at a temp. of .apprx.400° after which heating the pyridine insol. content was 57%, thus the pitch had a mesophase content of .apprx.57%. Fibers of 15 µm were spun at 390° in a N atm. and part were heated for 1 hr in an air-atm. furnace to 275° and held at this temp. for an addnl. hr to heat harden. The hardened fibers (.apprx.300g) were cut in 2.54 cm lengths, placed in a Pyrex beaker and in a sagger slowly (60% hr) heated to 500°, held at this temp. for 3 hr and then cooled to room temp. The fibers were then heated to 1000° in a graphite crucible and for 5 hr held at this 1000°. The cooled fibers were powdered (.apprx.200 µm) and mixed with coal tar pitch (100 parts powder to 80 parts pitch) and this mixt. was extruded at a pressure of 7-14 kg/cm<sup>2</sup> to a rod 2cm in diam. The rod was heated to 1000° at 60°/hr, held-at this temp. for 2 hr then in .apprx.1 hr heated to 3000° and there held for 2 hr. The rod had a coeff. of thermal expansion of 0.67 + 10-6/°C.

ST graphite electrode thermal expansion

IT Pitch



(graphite electrodes, with low thermal expansion)

IT Electrodes  
(graphite, from coal tar pitch, with low thermal expansion)

IT 7782-42-5, uses and miscellaneous  
RL: USES (Uses)  
(electrodes, from coal tar pitch, with low thermal expansion)

L35 ANSWER 52 OF 53 JAPIO (C) 2005 JPO on STN  
AN 2004-127913 JAPIO  
TI LITHIUM SECONDARY BATTERY  
IN KATO FUMIO; OURA TAKAFUMI; FUKUMOTO YUSUKE; YAMAMOTO NORIHIRO;  
FUJIWARA SHOZO  
PA MATSUSHITA ELECTRIC IND CO LTD  
PI JP 2004127913 A 20040422 Heisei  
AI JP 2003-160969 (JP2003160969 Heisei) 20030605  
PRAI JP 2002-223862 20020731  
SO PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined Applications, Vol.  
2004

IC ICM H01M004-02  
ICS H01M002-02; H01M004-58; H01M004-62; H01M004-66; H01M010-40

AB PROBLEM TO BE SOLVED: To improve charge and discharge cycle characteristics of a high energy-density lithium secondary battery, and improve or maintain discharge rate characteristics, low-temperature discharge characteristics, and safety (heat resistance).  
SOLUTION: The lithium secondary battery is prepared by using a negative electrode wherein active material composed of a mixture consisting of an artificial graphite particle A that has been obtained by kneading and granulating the base material made by crushing a bulk mesophase pitch, a pitch, and/or a thermosetting resin in a softened state and by making it carbonized/graphitized, and a spherical graphite particle B of which circularity is large, is fixed on the copper core material. This enables to improve the charge and discharge cycle characteristics of the high energy-density lithium secondary battery, and at the same time to provide the battery that is superior in the discharge rate characteristics, the low-temperature discharge characteristics, and safety (heat resistance).  
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L35 ANSWER 53 OF 53 JAPIO (C) 2005 JPO on STN  
AN 2000-156226 JAPIO  
TI NEGATIVE ELECTRODE FOR BATTERY AND NONAQUEOUS ELECTROLYTE SECONDARY BATTERY  
IN KOBAYASHI KOTARO; KOJIMA AKIRA  
PA SHIN KOBE ELECTRIC MACH CO LTD  
PI JP 2000156226 A 20000606 Heisei  
AI JP 1998-327923 (JP10327923 Heisei) 19981118  
PRAI JP 1998-327923 19981118  
SO PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined Applications, Vol.  
2000

IC ICM H01M004-58  
ICS H01M004-02; H01M004-62; H01M010-40

AB PROBLEM TO BE SOLVED: To prevent the delamination and cracking of a mixture layer while keeping high capacity without increasing a quantity of binding agent which inhibits the charging and discharging reaction, in a case when a rolled copper foil capable of providing high capacity and the lumped graphite powder easily causing the delamination of the mixture layer layer, are used in the negative electrode active material.  
SOLUTION: In a negative electrode for nonaqueous

*Applicant*

electrolyte secondary battery comprising a mixture including a carbon material and a polyvinylidene fluoride resin as a binder and a rolled copper foil, a mixture of lumped graphite powder and mesophase pitch graphite fibrous material is used as the carbon material. Whereby the density and adhesion of the mixture can be improved, and the irreversible capacity and the generation of gas at a high temperature can be inhibited.  
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